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Final Report

THE DEVELOPMENT OF  
A NONCRYOGENIC NITROGEN/OXYGEN  
SUPPLY TECHNIQUE

OFFICE OF PRIME RESPONSIBILITY

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by B. M. GREENOUGH

Prepared Under Contract NAS 9-10405

BIOTECHNOLOGY

LOCKHEED MISSILES & SPACE COMPANY

Sunnyvale, California

for

NATIONAL AERONAUTICS & SPACE ADMINISTRATION

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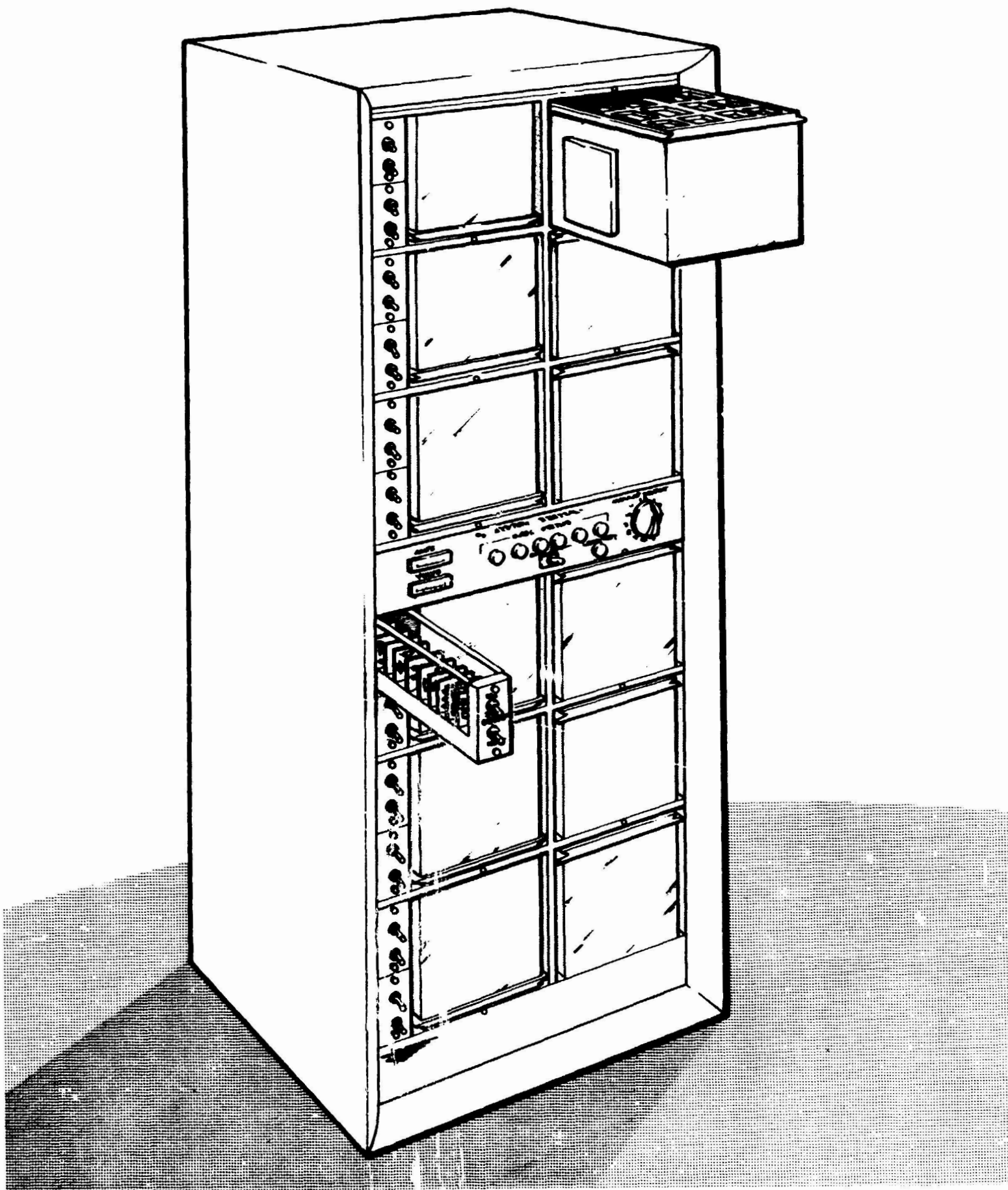
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**February 1971**

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Frontispiece - Space Station O<sub>2</sub>/N<sub>2</sub> Generation System

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### ABSTRACT

A development program was conducted in two phases to define the characteristics and requirements of an electrochemical oxygen/nitrogen supply technique for space station application. In Phase I, electrode formulations and structures suitable for use as anodes in an oxygen/nitrogen generator were experimentally investigated. A one-man model oxygen/nitrogen generator integrated with a space cabin atmosphere simulator was fabricated and successfully tested in Phase II. Data from these tests were used to update a computer routine model of the cabin atmosphere control using the oxygen/nitrogen generator technique. A specification and preliminary design for a 12-man oxygen/nitrogen generation system was prepared.

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## SUMMARY

A development program was conducted concerned with the use of a hydrazine/water electrolysis technique to provide both the metabolic oxygen for crew needs and the oxygen and nitrogen for cabin leakage makeup. A laboratory breadboard model one-man  $O_2/N_2$  generation system was integrated with a cabin and metabolic/leakage simulator to provide a testbed for determining cabin atmosphere control characteristics and for evaluating components.

Experimental electrodes were developed and were successfully operated in the one-man model system.

A zero-gravity closed reservoir system including a bubble separator was evaluated.

Control of the total pressure and oxygen partial pressure in the cabin simulator was demonstrated in a series of runs of the integrated system.

A computer routine model of the  $O_2/N_2$  system was updated and revised based on the experimental data from the integrated system testing. This updated computer model predicts adequate control of a 12-man space station under orbital conditions.

A preliminary design and specification for a full-scale 12-man  $O_2/N_2$  system suitable for space station use yielded a total system weight of 862 lb, including spares, and power consumption of 7,830 watts with redundancy and sparing to a reliability of 0.9986 for a 180-day mission.

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## Section 1 INTRODUCTION

For extended space-base and space-station manned missions, an oxygen/nitrogen cabin atmosphere will be utilized. The inert diluent will reduce the fire hazard of the oxygen and will enhance the physiological habitability of the environment.

Oxygen consumed metabolically by the crew is recovered from metabolic wastes in a water/waste management and regenerative life support system and is recycled to the cabin. Water electrolysis is a process considered for use in this cycle to recover oxygen from water and provide hydrogen for carbon dioxide reduction.

Losses in cabin atmosphere due to cabin leakage necessitate storage of oxygen and nitrogen for leakage makeup on long-duration missions. For a mission of less than 30 days, it may be practical to carry nitrogen and oxygen onboard the spacecraft using either cryogenic or high-pressure gaseous storage. For an extended mission, however, the weight penalty associated with cryogenic or high-pressure gaseous tankage is excessive.

The development program described herein is concerned with the use of a hydrazine/water electrolysis technique to provide both the metabolic oxygen for crew needs and the oxygen and nitrogen for cabin leakage makeup. With this system, oxygen and nitrogen are stored chemically as water and hydrazine in low-pressure (and therefore low-weight) tankage. This system also has the feature of providing automatic control of the space cabin total pressure and oxygen partial pressure.

The primary objectives of the program were to acquire test data on a laboratory model of the oxygen/nitrogen system and to provide a preliminary design for a 12-man prototype system. The program involved two major areas of effort consisting of (1) electrode and cell development and (2) the assembly and test of a one-man model system to

provide data for the preliminary design. A computer routine model of the oxygen/nitrogen system control of a space cabin atmosphere, developed previously under Contract NAS1-7706, was updated on the basis of the model system test data.

The specific tasks that were completed in meeting the program objectives were as follows:

- Electrode active material development
- Electrode structure and fabrication techniques
- Component selection
- Computer analyses
- Breadboard system assembly
- Breadboard system test
- Preliminary design of prototype system

These tasks were accomplished by analytical and experimental investigations in the following areas:

- Experimental electrodes were fabricated using various formulations, structures, and processing techniques. Performance was evaluated using a potentiostatic apparatus and an electrolysis cell test facility.
- Performance of system components was evaluated experimentally in a one-man model system. Performance characteristics were established for an oxygen partial pressure and total pressure control system, a cabin and metabolic/leak simulator, a zero-gravity water feed system, a zero-gravity bubble separator, and a hydrazine flow control system.
- Automatic control of a cabin simulator total pressure and oxygen partial pressure was demonstrated with the model oxygen/nitrogen generation system in a series of tests.
- Computer analyses were made in conjunction with the system testing to update the computer routine model. Test results were used to verify the updated computer routine.

- A preliminary system design effort culminated in the design of a 12-man oxygen/nitrogen generation system capable of providing a nominal supply of 26.1 lb/day of oxygen and 8 lb/day of nitrogen. The design includes a maximum load capability of 33.2 lb/day of oxygen and 22.6 lb/day of nitrogen.

The sections that follow in this report are concerned primarily with the areas of electrode development, system testing, and computer analyses. The computer routine and the preliminary system design have been documented separately with a computer utilization manual (Ref. 1) and preliminary design and specification report (Ref. 2).

## **Section 2**

### **RESULTS AND TECHNICAL DISCUSSION**

#### **2.1 ELECTRODE DEVELOPMENT**

##### **2.1.1 Electrode Design Requirements**

The electrode, an integral part of the electrolysis unit, must meet rigid electrical and physical requirements. (Refs. 3 and 4.)

The basic components of the electrode assembly in the LMSC electrolysis unit are: (a) rim, (b) catalytic surface, and (c) catalyst. Functions of these components are:

- The rim besides serving as a rigid support, provides a path for current and voltage.
- The catalytic surface is the center of the electrolysis process.
- The catalyst's primary function is to reduce the chemical activation overpotential.

The design of the rim and the use of platinum black in a Teflon catalytic surface were established at the beginning of the program. Accordingly, all developmental effort was centered on the fabrication and application of a Teflon/platinum (Pt) surface on a nickel screen.

Theoretically, a catalytic surface can be divided into two main parts, one to provide physical strength and "flow through" properties for reaction gases and reactants, and a second part to provide maximum reactive area for the electrolysis process.

In actual operation, the matrix must provide a wettable or hydrophilic surface for the electrolyte to function and a pore structure with enough hydrophobicity to transfer the

reaction gases and repel the electrolyte. The pore structure must not be too dense so as to block the gas flow.

Another property that is important is electrical conductivity. Resistance must be low. Thus, the ideal electrode might be one with a maximum hydrophilic conductive reaction area as well as a porous hydrophobic structure for efficient gas passage.

Attainment of the above with Teflon and platinum is primarily a fabrication problem. In order to improve application of experimental mixes and conductivity of the final matrix, acetylinic carbon was added. The carbon also helped reduce the hydrophobic properties of the Teflon.

#### 2.1.2 Test Facilities

The following two test facilities were available:

- Immersed Electrode Polarization Apparatus
- Laboratory Model Electrolysis Cell Apparatus

Immersed Electrode Polarization Apparatus. In order to study initial performance characteristics of the experimental electrodes, the Immersed Electrode Polarization Apparatus was designed and built. (See Fig. 2-1.) This apparatus is essentially two H cells – electrolysis and reference units, respectively – connected by an electrolyte bridge. The reference unit is equipped with three standard platinum electrodes, two for hydrogen and oxygen production and the third for a point of reference. The counter and working (experimental) electrodes are located in the electrolysis unit. The complete cell holds 300 cc of 30-percent potassium hydroxide solution.

A WENKING potentiostat is used to control and measure variations in voltage and current between the reference and working electrodes. Both units are vented to the hood.

This apparatus provides a quick and accurate evaluation of the effective surface area or reaction zone of candidate electrodes prior to final testing in the laboratory model electrolysis cell.

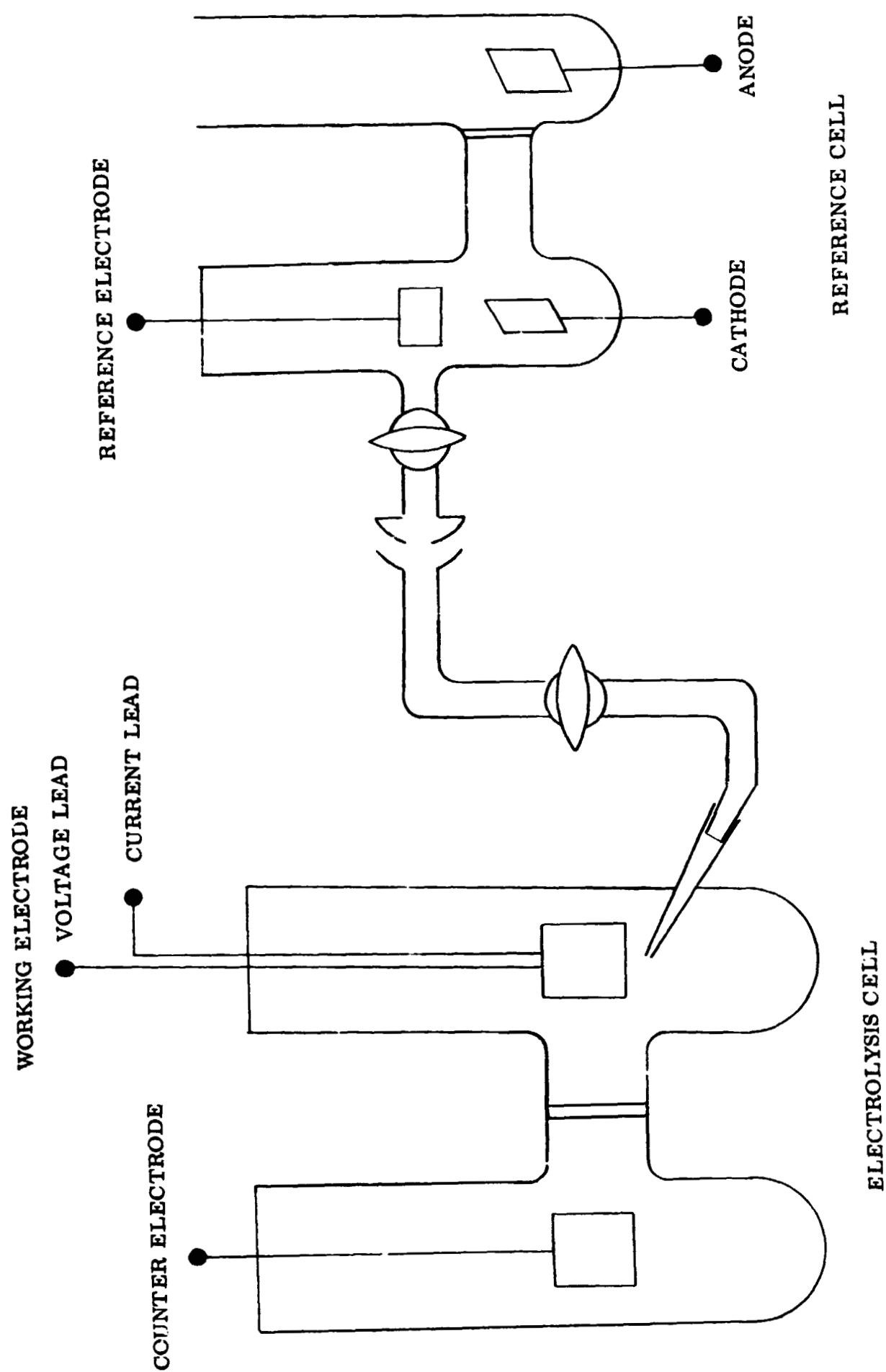


Fig. 2-1 Immersed Electrode Polarization Apparatus



Electrode samples were cut to a  $1 \times 2$  cm size and spotwelded on a nickel lead. Data points were obtained from the electrode samples by measuring the reference voltage vs. the working voltage for different current densities.

Laboratory Model Electrolysis Cell Apparatus. The Laboratory Model Electrolysis Cell Test Facility designed and built by LMSC (Contract NAS1-7706) is routinely used for the purpose of obtaining long-term electrolysis cell data and for general cell development. In contrast to the Immersed Electrode Polarization Apparatus, this test facility simulates actual operational conditions. Four test stations are available. Each station is provided with the necessary inputs of current-controlled electrolysis power, an auxiliary ac power source, coolant, feed water, and inert gas purge. Ammeters are used for current readout, and both digital and strip chart readouts are used for voltage. A complete description follows.

Two test stations are shown in Fig. 2-2. The electrolyte circulation loop components can be seen in this figure. Electrolyte leaving the electrolysis cell is discharged into a reservoir. A magnetically-driven, plastic centrifugal pump is used to circulate the electrolyte. From the pump, the electrolyte is passed through an all-plastic shell and tube heat exchanger in which cold ethylene glycol flows through the shell side. A plastic valve is used to adjust the electrolyte flow as indicated by a flowmeter. After passing through the flowmeter, the electrolyte is returned to the cell. (See Section 2.2.1 for a complete description of the cell configuration.)

After startup, the operation of a cell is completely automatic. The main control functions are cell temperature and water feed. Cell temperature control is maintained with a miniature thermostatic switch located at the electrolyte discharge from the cell. This switch provides a signal to a solenoid valve to provide coolant on demand. A heating circuit also is provided to permit testing at temperatures above ambient. Water feed control is achieved with a level control switch that provides a signal for liquid water feed to the reservoir on demand. Because the test stations were designed for automatic operation and unattended operation on a 24-hr a day basis, a number of safety

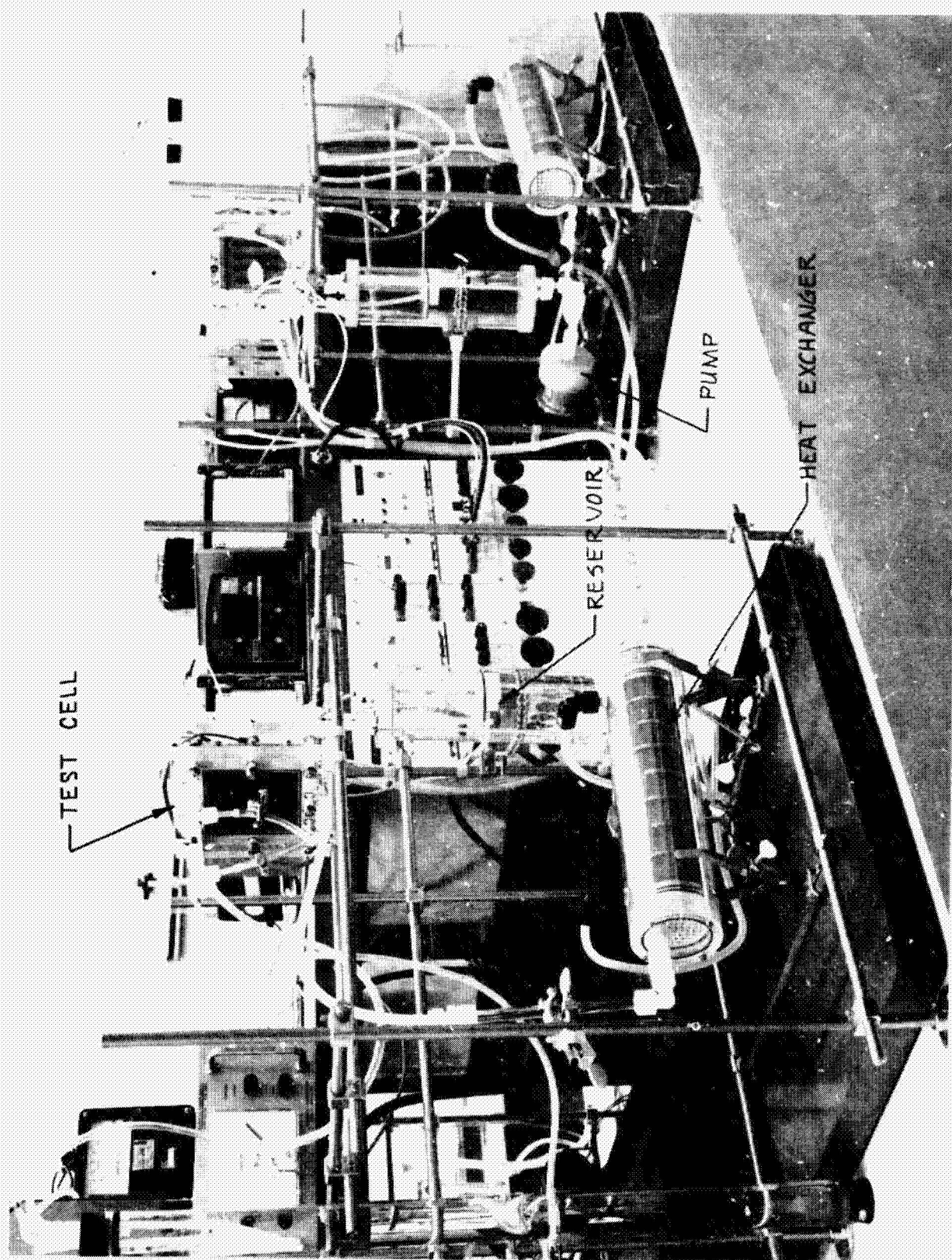


Fig. 2-2 Electrolysis Cell Test Stations

functions were built into the control panels shown in Fig. 2-3. During normal operation, panel lights indicate the electrolyte cooling and heating cycles, that the pump is operating, and that the purge-gas ( $N_2$ ) solenoids are closed. Unsafe operating conditions, if they occur, are indicated in the upper row of panel lights. Unsafe conditions would exist if there were no electrolyte flow; the electrolyte level in the reservoir were too high or too low; or the cell temperature were too high.

In the event of an unsafe condition, the safety circuit is activated to automatically shut down the test station. Electrolysis cell power is turned off; the pump, coolant, and water feed go off; and nitrogen purge to both sides of the cell is turned on. The panel light for whichever safety function shuts down the station remains on, indicating the cause of shutdown, until it is reset. All of the safety circuits contain latching relays to prevent the station from being reactivated automatically before the cause of the unsafe condition can be rectified.

Differential pressure control between the gas and electrolyte phases is accomplished by discharging electrolyte from the cell below the level of the inlet. This provides a suction pressure in the electrolyte so that the generated gases can be discharged at ambient pressure and still maintain a positive gas-over-liquid pressure. Although it is not necessary for maintaining sufficient gas-back pressure, the generated gases are generally discharged into bubblers with a few inches of water back-pressure to give a visual indication of gas generation.

Cell performance data are automatically recorded on a continuous basis with an automatic data logging system utilizing a high-speed typewriter, digital clock, digital voltmeter, and multichannel scanner.

### 2.1.3 Cell Tests - Operating Parameters

Voltage vs. Current Density. The voltage-current density relationship (polarization curve) for an electrolysis cell is the primary descriptor of its performance. Because the gas generation rate is Faradaic, the power required to produce a given amount of

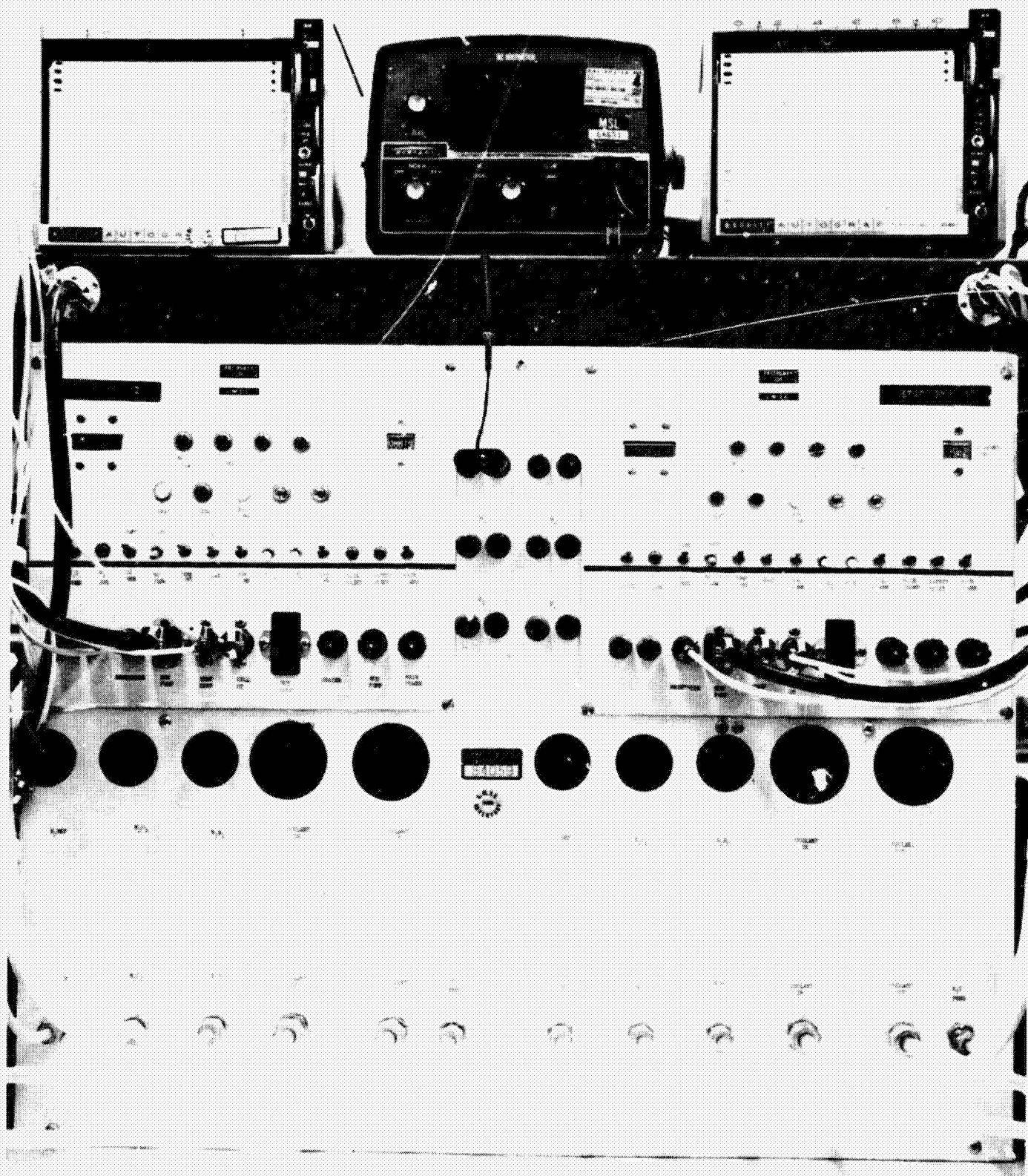


Fig. 2-3 Electrolysis Test Station Control Panels

gas is directly proportional to the cell voltage. It is, therefore, desirable to have a cell with a "flat" polarization curve, i.e., minimum voltage change with increasing current. This also applies to the small polarization apparatus measurements.

The cell voltage is the sum of four components:

$$V_{\text{cell}} = V_{\text{oc}} + V_{\text{ir}} + \eta_{\text{O}_2} + \eta_{\text{H}_2} \quad (2.1)$$

where

$V_{\text{oc}}$  = theoretical open-circuit cell potential

$V_{\text{ir}}$  = IR loss

$\eta_{\text{O}_2}$  = oxygen overvoltage

$\eta_{\text{H}_2}$  = hydrogen overvoltage

The theoretical open-circuit cell potential for hydrogen and oxygen is 1.23 V. However, in an actual device, the open-circuit potential is closer to 1.0 V due to a mixed potential at the oxygen electrode. When current is passed between the electrodes, a voltage drop (or IR loss) is produced because of the cell resistance. For a given cell configuration, the resistance is fixed and the IR loss is proportional to the applied current.

The overvoltage contributions to the cell voltage are a complex function of catalytic activity of the active electrode material and the mechanism of the reaction at each electrode. A number of contributions to the literature have advanced theories to explain the existence of overvoltages and mechanisms of the reactions. These theories have been conflicting in most cases and no one explanation has gained universal acceptance.

While it is possible to determine electrode overvoltages by physical measurement in bulk electrolyte (Ref. 5), the use of these data to predict electrode performance in a practical cell is complicated by the presence of the matrices. A more practical

approach that was taken in this program was to insert a standard hydrogen reference electrode in the external electrolyte circulation loop. The voltages that were then monitored were related by:

$$V_{\text{cell}} = V_{\text{O}_2} + V_{\text{H}_2} \quad (2.2)$$

where

$V_{\text{O}_2}$  = oxygen electrode potential vs. standard hydrogen

$V_{\text{H}_2}$  = hydrogen electrode potential vs. standard hydrogen

Comparing Eq. (2.2) with Eq. (2.1) it is seen that  $V_{\text{O}_2}$  and  $V_{\text{H}_2}$  each include a part of the open-circuit potential,  $V_{\text{OC}}$ , and the IR loss,  $V_{\text{IR}}$ , in addition to the over-voltage. At open circuit,  $V_{\text{H}_2}$  is zero and  $V_{\text{OC}}$  is equal to  $V_{\text{O}_2}$ . Since only the total cell resistance and not the resistance distribution across the cell was measured, the IR loss contribution to each electrode and absolute values of overvoltage could not be determined from the measurements of  $V_{\text{O}_2}$  and  $V_{\text{H}_2}$ .

These potentials, which will be referred to in subsequent sections as "electrode polarization," are useful parameters for evaluating electrolysis cell performance and give additional insight into the operating characteristics of different electrode and matrix materials and the effect of other operating parameters.

A typical electrolysis cell polarization diagram is shown in Fig. 2-4. The voltages shown are initial values and do not reflect long-term performance changes. The polarization of the oxygen electrode is more than three times as severe as that of the hydrogen electrode, and it is evident that cell performance improvement is more likely to be achieved by reducing the oxygen electrode polarization.

Temperature and Electrolyte Concentration. The cell temperature and KOH electrolyte concentration are operating parameters that are independent of each other in that each



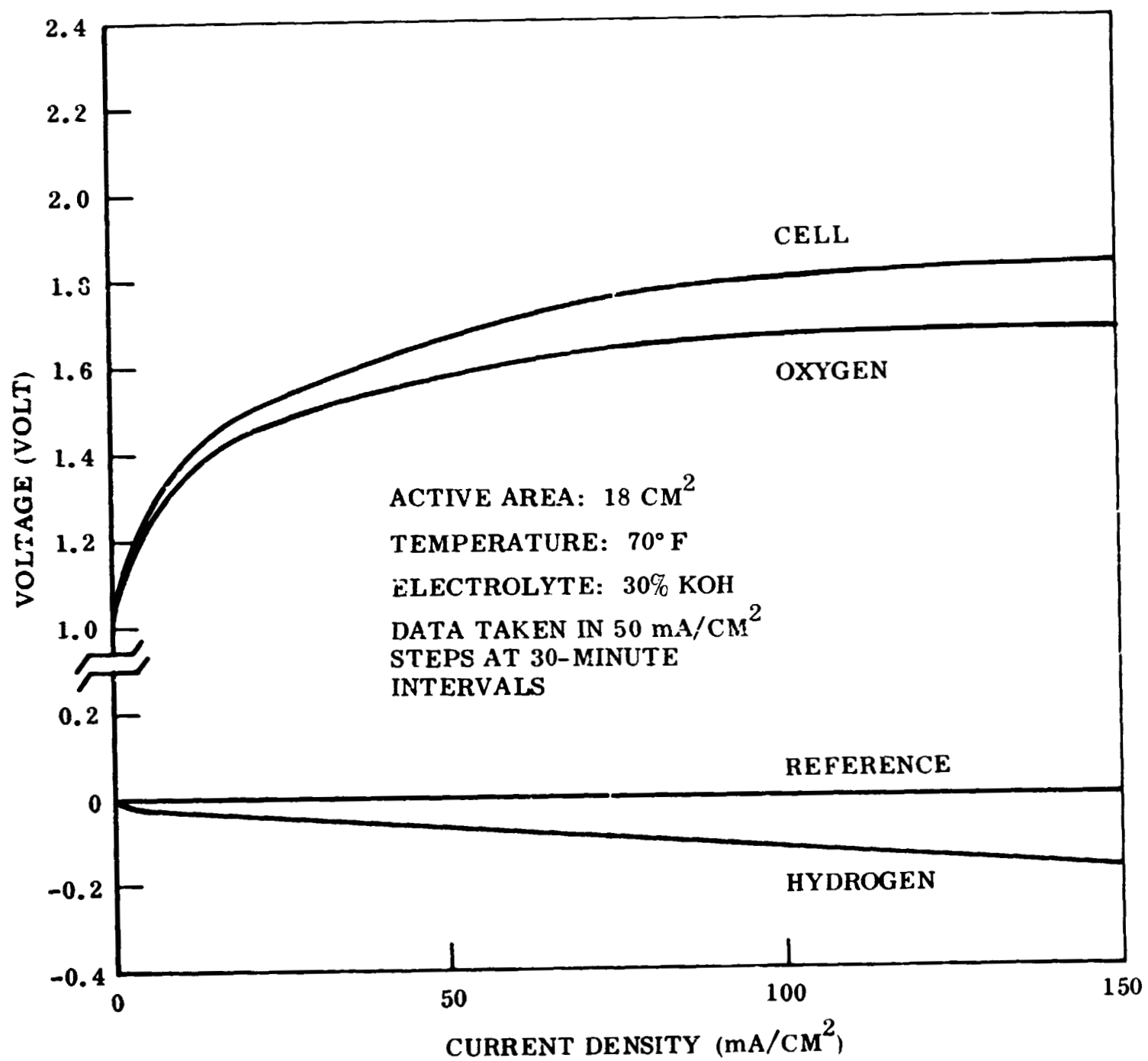


Fig. 2-4 Electrolysis Cell Polarization

can be set arbitrarily and controlled without regard to the other. Their effects on the operating characteristics of the cell, however, are interrelated. The first criterion for choosing the optimum electrolyte concentration is to minimize the internal cell IR loss. As can be seen in Fig. 2-5, the minimum electrolyte resistance occurs between 25 and 30 weight percent.

The effects of temperature are two-fold: first, the temperature affects the operating cell voltage; and second, it affects the absolute humidity of the generated gases.

#### 2.1.4 Experimental Results and Discussion

A summary of the experimental electrode testing that was conducted is presented in Table 2-1.

The following materials were used in the experimental work:

<u>Material</u>	<u>Source</u>
Teflon (TFE) Powder TL 126	Liquid Nitrogen Processing Corp.
Teflon Industrial Finish	E. I. Dupont DeNemours & Co.
50% TFE Teflon Suspension	
Black Platinum	Englehard Co.
Nickel Powder	LMSC
Graphite	—
Expanded Nickel Screen	Exmet Diemesh Corp.

##### 2.1.4.1 Preliminary Formulations

Initial work was confined to a Teflon/catalyst mix with the following desired properties:

- It must be easily applied to a nickel screen.
- The final catalytic surface must be hydrophilic, conductive, and porous.
- It must adhere to the screen.



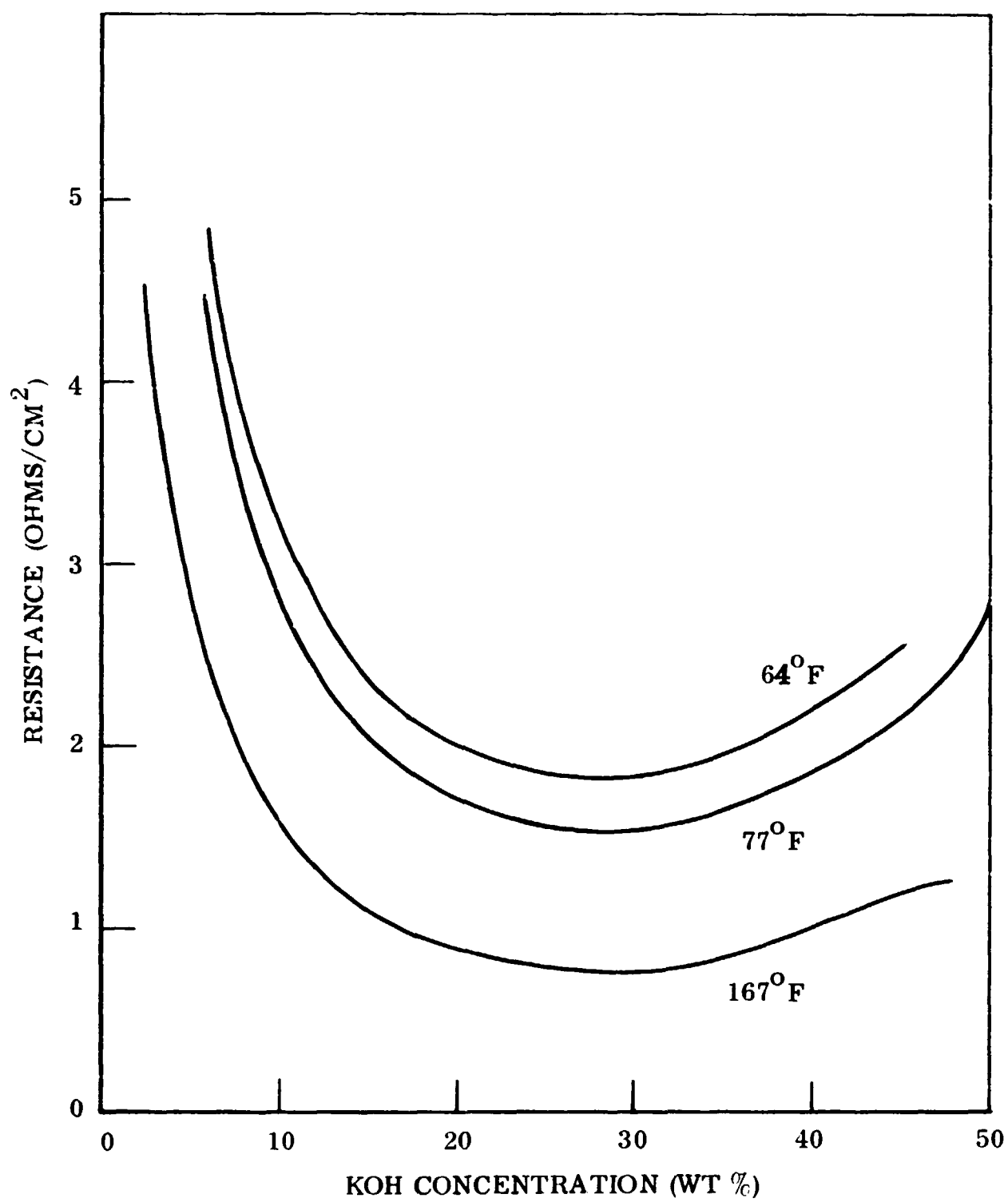


Fig. 2-5 Electrolyte Resistance vs. Concentration

Table 2-1  
EXPERIMENTAL ELECTRODE TEST SUMMARY

Electrode Number	Catalyst Load (mg/cm <sup>2</sup> )	Type of Test	Anode Voltage (V)	Current Density (mA/cm <sup>2</sup> )	Test Time (hr)	Comments
T-1	Ni 87	Conductivity				Electrode surface nonconductive
T-2	Ni 50	Conductivity				Electrode surface nonconductive
T-3	Ni	Processing				Mixture would not adhere to substrate
T-4	Pt 13	Processing				Formed nonconductive film
T-5	Pt 12	Processing				Formed nonconductive film
T-6	Pt 9	Processing				Formed nonconductive film
T-7 through T-18	Pt	Processing				No measurements made
T-19	Pt 20	Polarization	1.85	75	2	Disintegrated
T-20	Ni 20	Polarization				Immersed test - see Fig. 2-6
T-21	Ni 20	Polarization	2.00	50	2	Immersed electrode polarization test
T-22	Ni 30	Polarization	2.00	125	2	Immersed electrode polarization test
T-23	Ni 30	Polarization	2.00	150	2	Immersed electrode polarization test
T-24	Ni 45	Polarization	2.00	150	2	Immersed electrode polarization test
T-25	Pt	Processing				Mixture did not adhere to substrate
T-26 through T-28	Pt	Processing				
T-29	Pt 16	Polarization	2.00	50	2	Immersed electrode polarization test
T-30	Pt 20	Polarization	2.00	38	2	Immersed electrode polarization test
T-31	Pt 20	Polarization	2.00	80	2	Immersed electrode polarization test
T-32	Pt 40	Polarization	2.00	35	2	Immersed electrode polarization test
T-33	Pt 80	Polarization	2.00	115	2	Immersed electrode polarization test
T-34	Pt 25	Processing				No measurements made
T-35	Pt 26	Processing				No measurements made
T-36	Pt 25	Processing				No measurements made
T-37	Pt 26	Processing				No measurements made
T-38	Pt 20	Processing				No measurements made
T-39	Pt 21	Processing				No measurements made
T-40	Pt 25	Polarization	1.98	150	2	Immersed electrode polarization test
T-41	Pt 25	Polarization	2.00	150	2	Immersed electrode polarization test
T-42	Pt 33	Polarization	2.00	150	2	Immersed electrode polarization test
T-43	Pt 22	Processing				No measurements made
T-44	Pt 16	Polarization	1.82	150	2	Immersed electrode polarization test
T-45 through T-52	Pt	Processing				No measurements made
T-53	Pt 30	Polarization	1.82	150	1	Immersed test - see Fig. 2-7
T-54	Pt 30	Polarization	1.80	150	1	Immersed test - see Fig. 2-7
T-55 through T-62	Pt	Processing				No measurements made
T-63	Pt 35	Polarization	2.2	150	7	Immersed electrode polarization test
T-64	Pt 35	Polarization	1.8	120	1	Immersed test; cellulose acetate binder
T-65 through T-70	Pt	Processing				Scale-up to 18 cm <sup>2</sup> size
T-71	Pt 22	Electrolysis	2.18	150	294	See Fig. 2-10, 18 cm <sup>2</sup>
T-72	Pt 22	Electrolysis	2.10	150	200	See Fig. 2-10, 18 cm <sup>2</sup>
T-73 through T-86	Pt	Processing				Scale-up to 90 cm <sup>2</sup> size
T-87	Pt 20	Electrolysis	2.3	150	<1	
T-88	Pt 6	Electrolysis	2.4	150	1	
T-89	Pt 11	Electrolysis	2.3	150	2	
T-90 and T-91	Pt 12	Processing				No measurements made
T-92	Pt 20	Electrolysis	2.5	150	7	
T-93 through T-114	Pt	Processing				No measurements made
T-115	Pt 20	Electrolysis	2.3	150	7	
T-116	Pt 20	Electrolysis	2.3	150	3	
T-117 and T-118	Pt	Processing				No measurements made
T-119	Pt 20	Electrolysis				See Fig. 2-10
T-120	Pt 22	Electrolysis	2.3	150	6	
T-121	Pt	Processing				No measurements made
T-122	Pt 22	Electrolysis	2.10	150	136	See Fig. 2-11
T-123	Pt 22	Electrolysis	2.16	150	142	See Fig. 2-11
T-124	Pt 22	Electrolysis	1.88	150	135	See Figs. 2-11 and 2-12
T-125	Pt 22	Electrolysis	1.80	150	35	See Figs. 2-11 and 2-13
T-126	Pt 22	Electrolysis	2.10	150	70	See Figs. 2-11 and 2-14
T-127	Pt 22	Structure				See Fig. 2-8

Accordingly, many experimental formulations were prepared using various forms of Teflon and varying amounts of catalyst. Nickel powder and platinum black were used. The general process was to weigh the ingredients and to mix them with an easily removable solvent such as toluene. The resulting paste was spread on the expanded nickel screen, dried at room temperature, and pressed, either by rolling or pressing at a relatively high pressure. In some cases, the electrode was heat treated.

The initial electrodes did not meet all of the desired properties. Excessive pressures rendered the Teflon a completely hydrophobic film, and excess Teflon had the same effect. Insufficient Teflon did not give the required structural strength to the electrode, and the catalytic surface disintegrated in the electrolyte. Excessive amounts of carbon tended to decrease the structural strength.

There were difficulties encountered in the initial application of the paste on the screen. The fluid paste tended to go through the screen. This problem was solved by backing the screen with a porous Teflon sheet prior to application of the paste. The sheet was then stripped off the completed electrode.

Later on, the nickel screen was coated with a microfilm of Teflon prior to application of the matrix.

The first series of electrodes was prepared using nickel as a catalyst rather than platinum in an effort to reduce experimental costs. These electrodes were relatively simple - merely a mix of Teflon, catalyst, and lubricant applied to a nickel screen. Electrode number T-20 is an example. Its formulation is as follows:

Teflon Powder (TL-126)	21.7 Percent
Nickel Powder	72.0 Percent
Graphite	6.3 Percent

The three ingredients were blended and approximately 0.1 gm of mineral oil was added. The resulting paste was rolled on a screen until a smooth catalytic surface

was formed. After heat treatment at 300°F for 20 minutes, the electrode was washed with toluene to remove the mineral oil, then dried at 100°C.

Conductivity and wettability of the electrode were satisfactory. However, it was not structurally strong. After several test cycles in the immersion polarization apparatus, the catalytic surface disintegrated when subjected to electrolytic action. (See Fig. 2-6.)

In order to improve the structural strength of the matrix, a commercially prepared Teflon suspension was used with the TL-126. Electrode number T-20A is an example.

Teflon Powder	24 Percent
Teflon Suspension	10 Percent
Nickel	48 Percent
Graphite	18 Percent

This electrode also disintegrated.

While certain techniques involved in the handling of Teflon mixes were developed in the work with nickel powder, it was found that substitution of platinum for nickel called for modification of procedures. Also, a decision was made to concentrate on using the Teflon suspension rather than the dry Teflon powder.

Formulations T-53 and T-54 were the first experimental platinum electrodes with electrical and physical characteristics approaching those of the acceptable commercial electrode. The formulation is as follows:

Teflon Suspension	40 Percent
Black Platinum	50 Percent
Graphite	10 Percent

The platinum and graphite were thoroughly blended in 3 to 5 cc of toluene. Then, the Teflon suspension was weighed into the mix and the formulation was worked with a spatula into a homogeneous paste. The paste was spread on a Teflon-backed screen

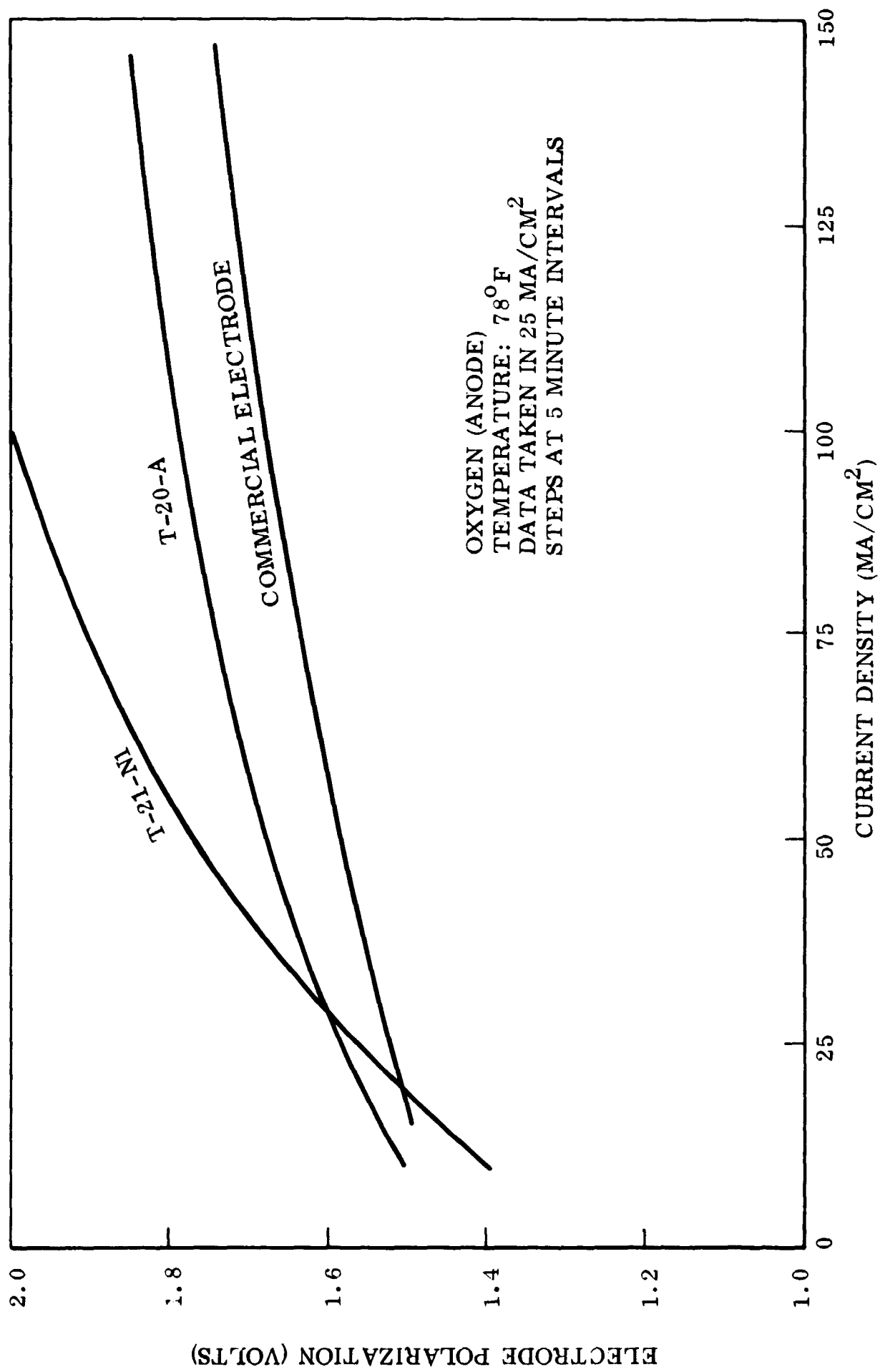


Fig. 2-6 Polarization of Experimental Nickel Electrodes

and rolled gently until it adhered to the screen. The electrode was dried (toluene removed) at room temperature for 2 hr, washed with distilled water, and finally dried in a 100°C oven.

Both T-53 and T-54 were porous, hydrophilic, conductive, and had sufficient structural strength to withstand the action of the 30-percent KOH solution. (See Fig. 2-7.)

During this phase of the work some effort was made to improve the porosity by incorporating a soluble material into the mix and, as a final step, washing the material out, leaving a porous matrix. This method was found to be generally unsatisfactory in that complete removal of the additive was difficult; hence, an impurity in the matrix.

Subsequent mixes using this basic formulation and varying the percentage of platinum from 40 to 60 percent showed that a satisfactory electrode could be prepared. It also was quite evident that the amount of pressing and rolling of the final catalytic surface had a marked effect upon the electrical uniformity of the final electrode. Excessive rolling tended to form a nonporous, nonconductive film. In general, the small electrodes were subjected to four passes through a small hand calendering device.

This phase of the work yielded an electrode that was conductive, hydrophobic, and offered enough catalytic area for efficient oxygen production in an immersed electrode apparatus.

Figures 2-8 and 2-9 are microphotographs of typical catalytic surfaces of an experimental electrode and a commercial electrode. Note the porosity.

#### 2.1.4.2 Electrodes for the Electrolysis Test Setup

Electrodes for this application were scaled up to 18 cm<sup>2</sup> and were circular. Different requirements presented new problems.

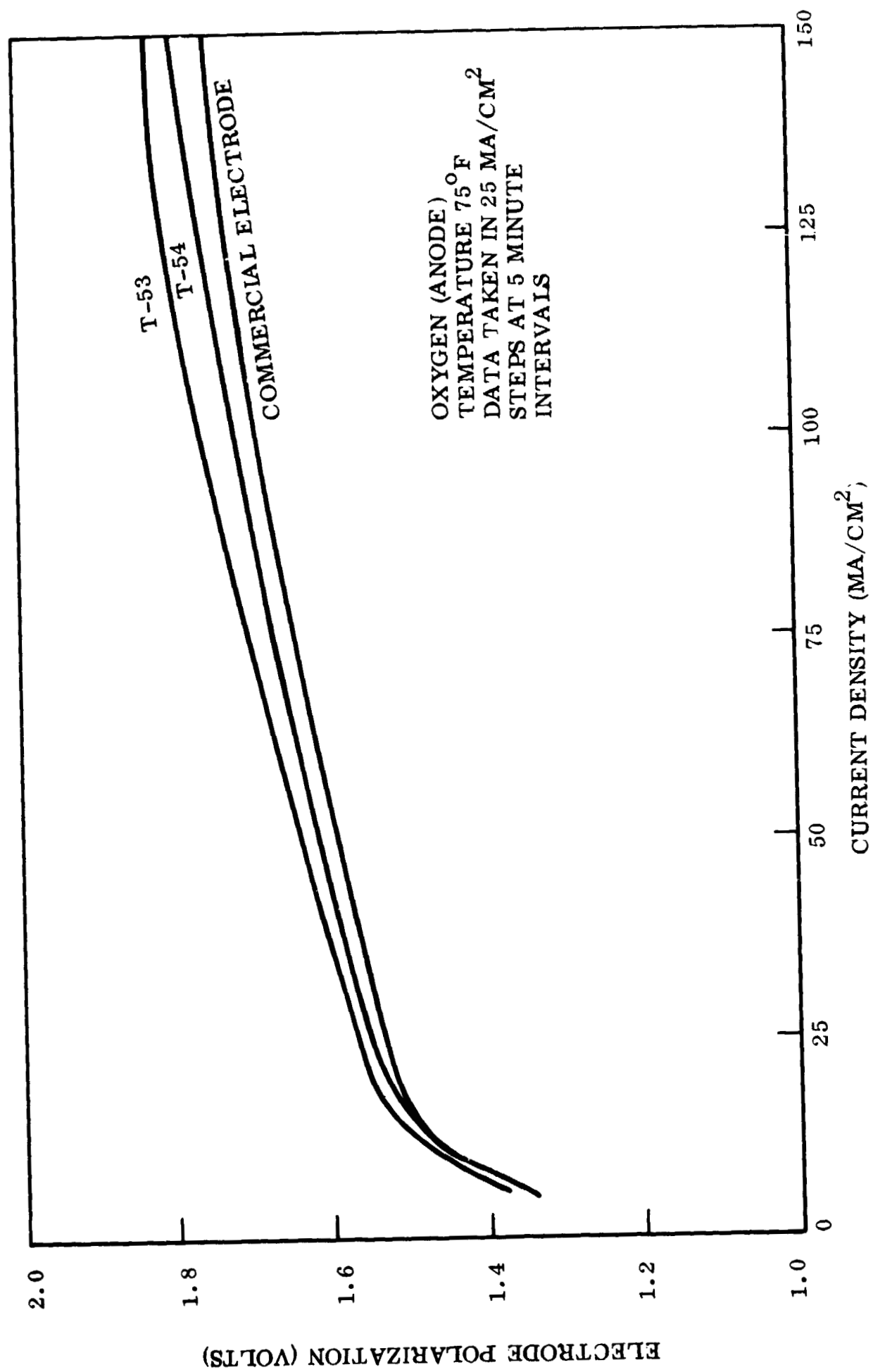


Fig. 2-7 Polarization of T-53 and T-54 Electrodes



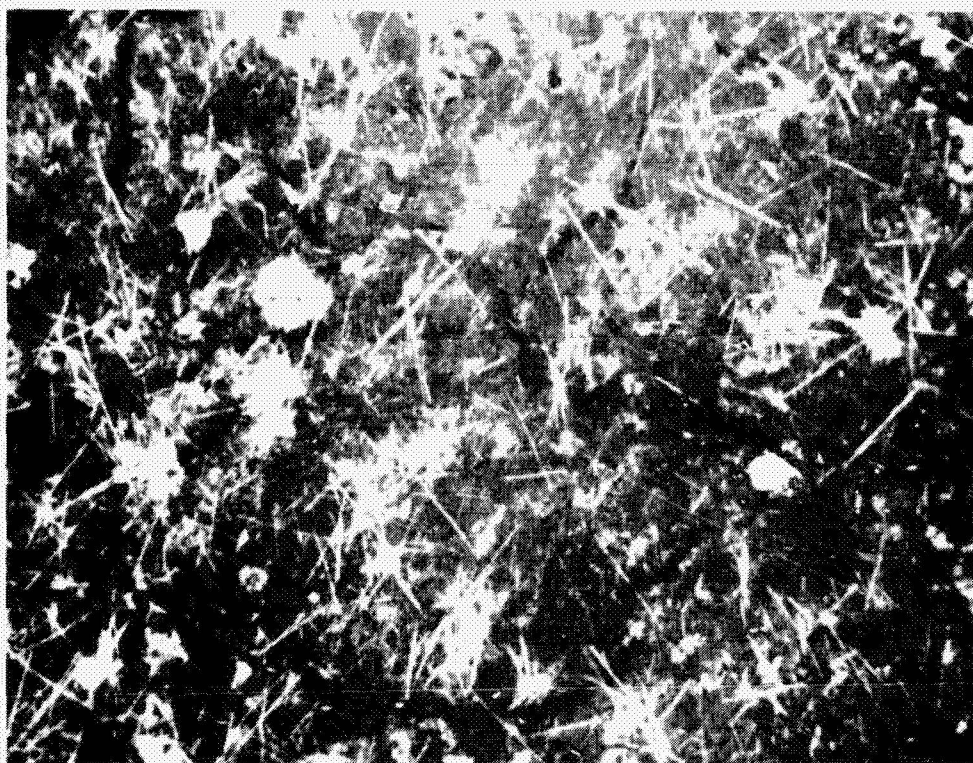


Fig. 2-8 Micrograph of T-127 (80 $\times$ )



Fig. 2-9 Micrograph of Commercial Electrode (80 $\times$ )



The two primary electrode requirements for efficient cell operation are as follows:

- The electrode must offer a maximum hydrophilic catalytic surface to the electrolyte.
- The porous inner electrode must be hydrophobic, so that the reaction gases will be transferred.

A totally wet electrode tends to offer a barrier to the gaseous reaction products, and the bubbles of gas, so blocked, will reduce the available catalytic surface of the electrode. A rapid rise in voltage is an indication of this condition. By cutting the power to the cell, the electrode will recover, but will again rise rapidly when power is restored.

A different approach was made at this point. A slightly hydrophobic carbon electrode was prepared first. Then a platinum/Teflon catalytic surface was applied to the carbon surface. T-71 is an example.

#### Carbon Electrode.

Acetylinic Carbon	58 Percent
Teflon Suspension	42 Percent

One cc of toluene was added to the above and the blend was mixed. The paste was spread on a Teflon-backed screen, dried at room temperature, and rolled until a porous, slightly hydrophobic film was obtained.

#### Platinum Electrode.

Black Platinum	50 Percent
Teflon Suspension	50 Percent

One cc of toluene was added to the above and, after mixing, the paste was spread on the carbon surface, dried at room temperature, and washed with distilled water. The

finished electrode was installed in the laboratory electrolysis unit. Results are shown on Fig. 2-10. The platinum loading was 22 mg Pt/cm<sup>2</sup>.

This electrode compared quite favorably with some of the better commercial electrodes. At the end of 294 hr, the cell voltage had risen from 2.04 to 2.39 V, and the oxygen voltage had risen from 1.6 to 2.18 V.

T-72 and T-119 (Fig. 2-10) are examples of electrodes prepared in this manner. Catalytic loads are 25 and 30 mg/cm<sup>2</sup> Pt, respectively.

Figure 2-11 shows the laboratory performance of experimental electrodes that were used later in the one-man unit discussed below. These electrodes were all made from the same formulation in the same manner.

#### 2.1.4.3 Large Electrodes for One-Man Unit

These electrodes were 90 cm<sup>2</sup>. Two methods of preparation evolved from the work.

- A catalyst/Teflon/carbon paste is prepared and applied to a screen.
- A carbon electrode is prepared first and a catalyst/Teflon matrix is applied to the surface.

Screen Preparation. The screen was given a dilute sulfuric acid wash followed by a distilled water rinse. It was found that a micro-coat of Teflon on the screen improved the gas flow through the matrix as well as increased the adhesion of matrix to screen.

Treatment of Matrix. Rolling, pressing, and temperatures were kept to a minimum in order to combat the film-forming tendencies of Teflon.

#### Usable Formulations.

Teflon Suspension	40 to 60 Percent
Platinum Black	55 to 45 Percent
Acetylinic Carbon	10 Percent

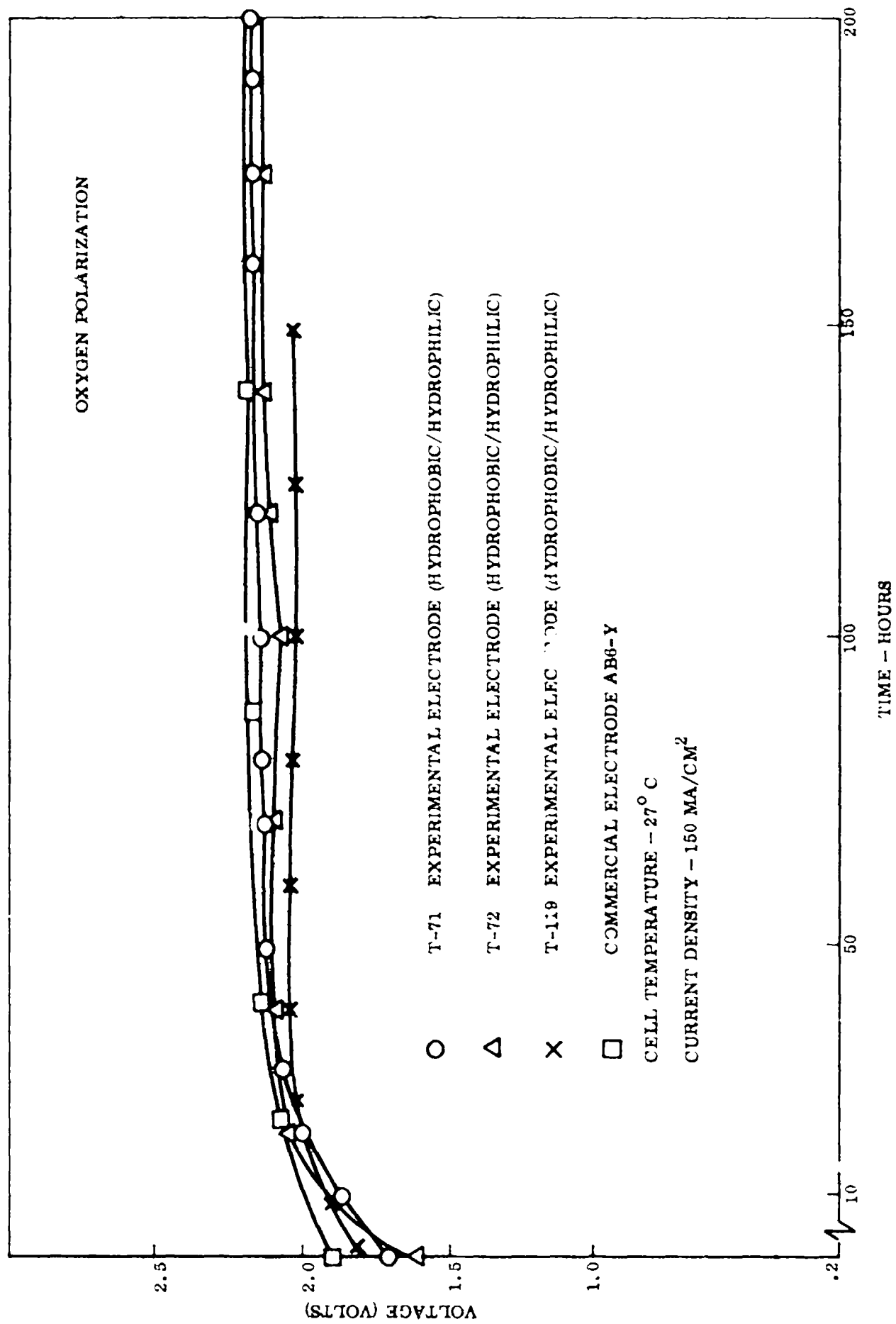


Fig. 2-10 Voltage vs. Time Curves for T-71, T-72, T-119, and a Commercial Electrode Assembly

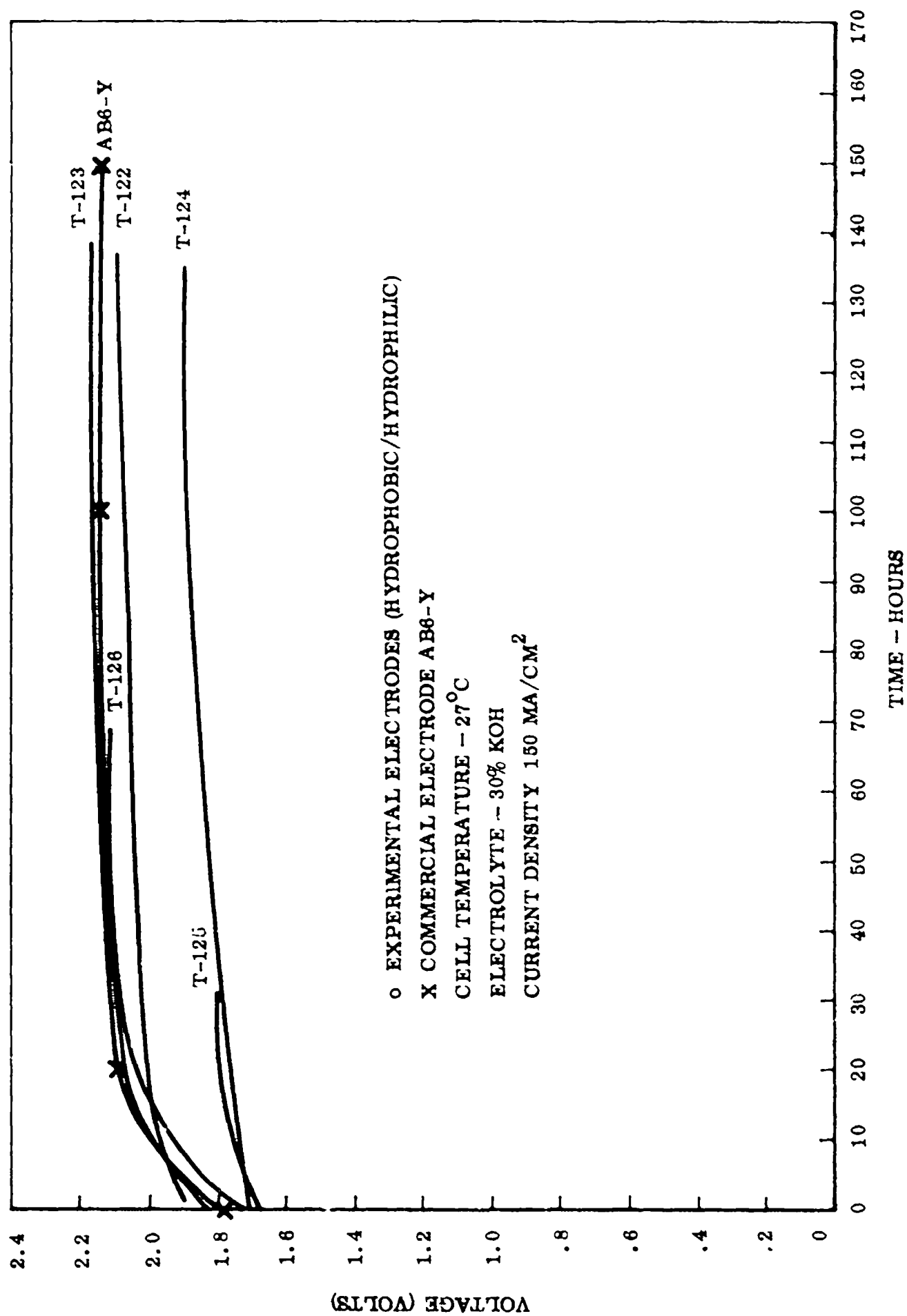


Fig. 2-11 Voltage vs. Time Curves for T-122 Through T-126

This mix, diluted slightly with toluene, can be applied to either a carbon/Teflon surface or a Teflon-backed screen. After drying at room temperature for 2 hr, the matrix can be rolled (calandered) four passes. It is a good idea to cover the surface with a thin polyethylene or wax paper sheet prior to rolling to prevent matrix pickup. Commercial wax paper was found to be satisfactory.

After the rolling operation, the matrix can be washed with toluene, dried at 100°C or less, and finally washed with distilled water.

Initial Performance Checks. Estimations of hydrophobic and hydrophilic properties can be made by dropping a drop of water on the surface. The water will remain a drop and will not wet a hydrophobic surface. A hydrophilic surface will wet, causing the drop of water to spread out. It is possible to make a rough estimation of contact angle and attending hydrophobicity or hydrophilicity of the catalytic surface.

The conductivity can be checked with a volt-ohmmeter. An interesting check on catalytic activity can be made by dropping one drop of methyl alcohol on a small portion of the electrode. A good electrode will cause the alcohol to burst into flame.

These various tests were used to provide a qualitative assessment of electrodes.

The final check is the installation of the electrode in the electrolysis unit. Figures 2-12, 2-13, and 2-14 show the electrical performance of three typical electrodes.

Analytical Work. Throughout the development period the purity of the oxygen generated by the experimental electrodes was monitored at various times by infrared and gas chromatographic analysis. No contaminants were detected.

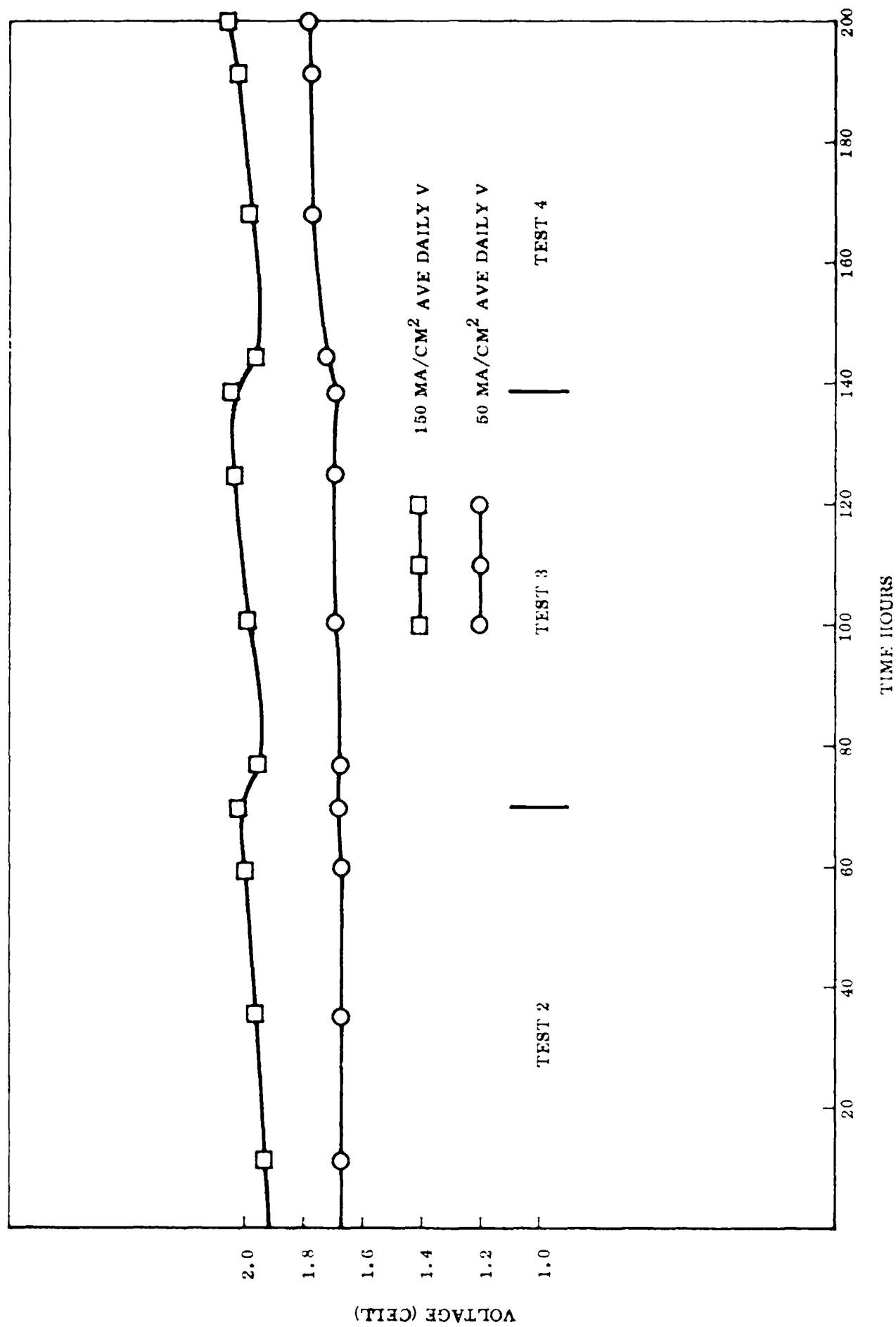


Fig. 2-12 T-124 Performance in O<sub>2</sub>/N<sub>2</sub> System

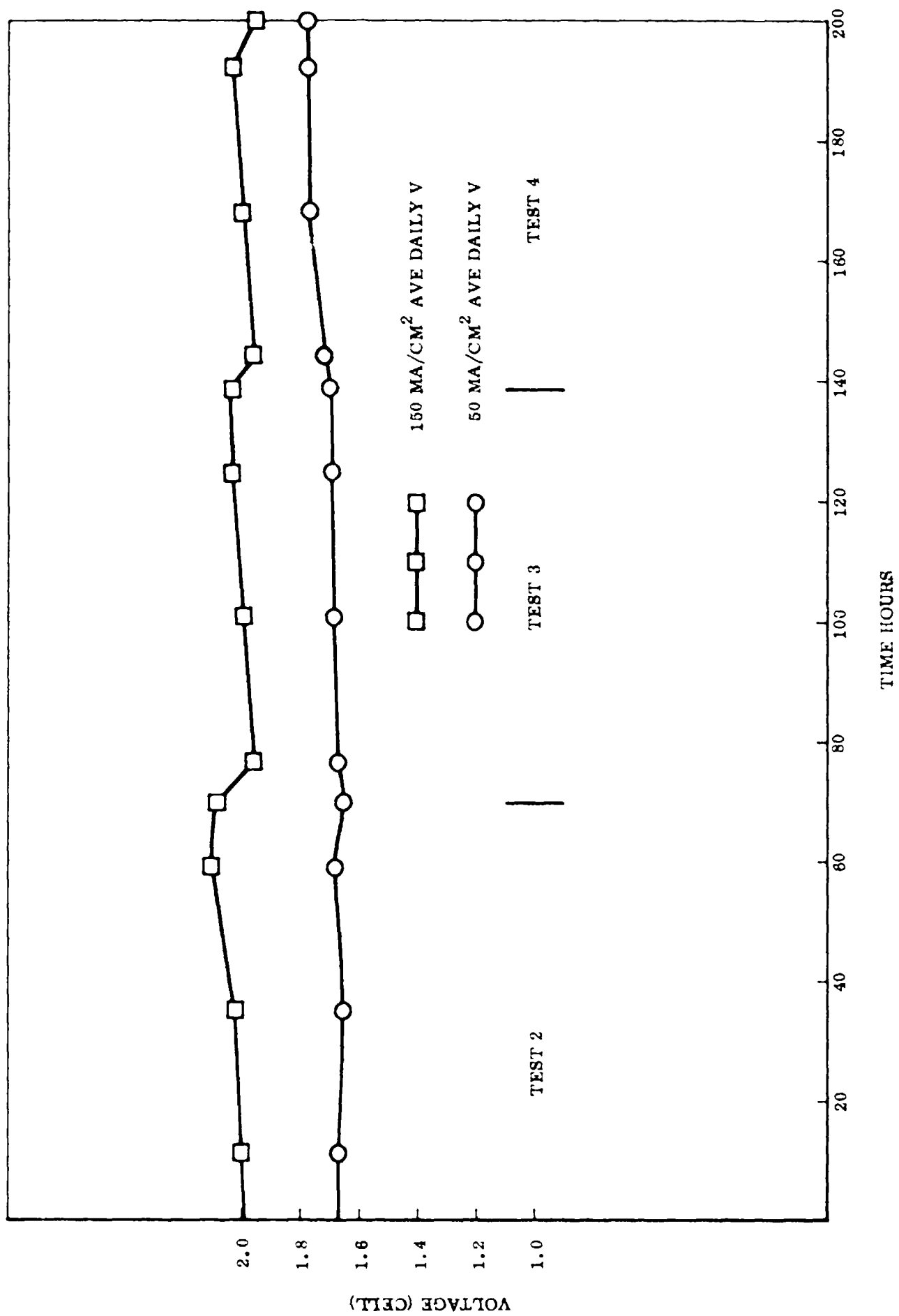


Fig. 2-13 T-125 Performance in O<sub>2</sub>/N<sub>2</sub> System

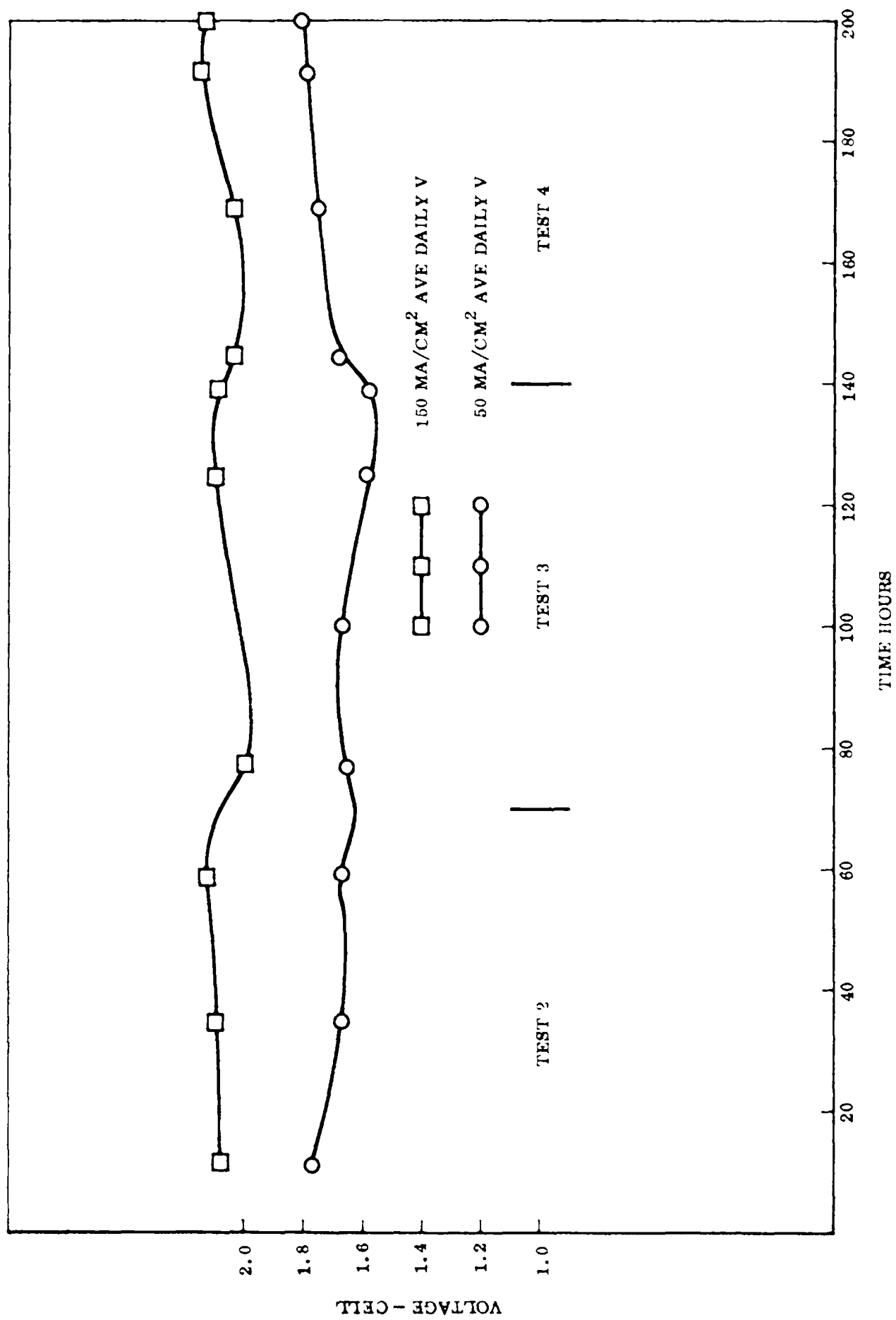


Fig. 2-14 T-126 Performance in O<sub>2</sub>/N<sub>2</sub> System



## 2.2 ONE-MAN MODEL SYSTEM DESCRIPTION

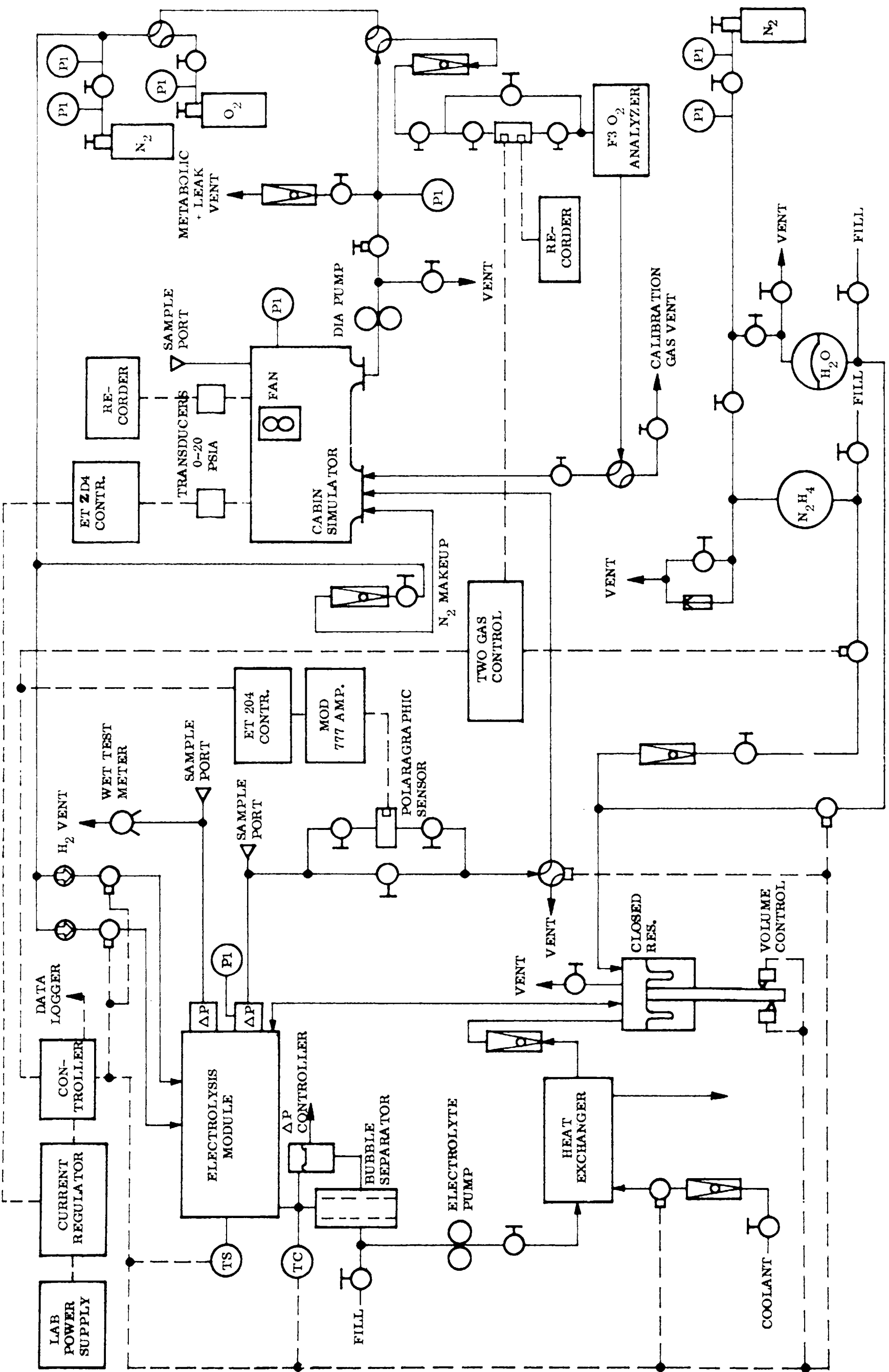
A laboratory breadboard hydrazine/water electrolysis system was assembled to provide a testbed for evaluating the oxygen/nitrogen generator technique. The gas generation system was sized to nominally provide a one-man metabolic supply and 1/12 of a space station cabin leakage makeup. This system was integrated with a cabin and metabolic/leak simulator and was instrumented to provide automatic control of the atmosphere in the cabin simulator. A schematic of this testbed is shown in Fig. 2-15.

### 2.2.1 Oxygen/Nitrogen Generation System

The oxygen/nitrogen generator consists of a closed-loop, liquid-electrolyte circulation system in which the gas generator, a stack of electrolysis cells, is operated in conjunction with accessories that provide for water and hydrazine feed, waste heat removal, differential pressure control, and bubble separation. Thirty percent potassium hydroxide is used as the electrolyte. Figure 2-16 shows the installed oxygen/nitrogen generation system. In this system, all surfaces in contact with the electrolyte, with the exceptions of the electrodes and matrix support screens, are composed of plastic.

Electrolysis Cell Stack. The electrolysis cell stack is comprised of 17 unit cells of the configuration shown in Fig. 2-17. The unit cell is designed so that liquid electrolyte flows through the center of the cell between two absorbent matrices contiguous to the operating electrodes. The matrices provide phase separation between the liquid in the center of the cell and the gases being generated on the open structure of the operating electrodes. Manifolds for the generated gases and the electrolyte are located on the periphery of the cell spacers; O-rings are used for sealing purposes. Cutaway views of the cell assembly are shown in Figs. 2-18 and 2-19.

The electrode assemblies consist of a rim, a spacer, support screens, an asbestos matrix, and an active electrode material. The rim is used for current take-off; a bent-over tab, which extends outside the cell, is used to interconnect electrodes in the stack of cells. The electrode spacer and support screen assembly is spotwelded to the rim and serves as a mechanical support and current distributor for the active material. The electrode consists of a mixture of black platinum catalyst and Teflon applied to a nickel screen substrate.



2-29

Fig. 2-15 One-Man Model  $O_2/N_2$  System Schematic





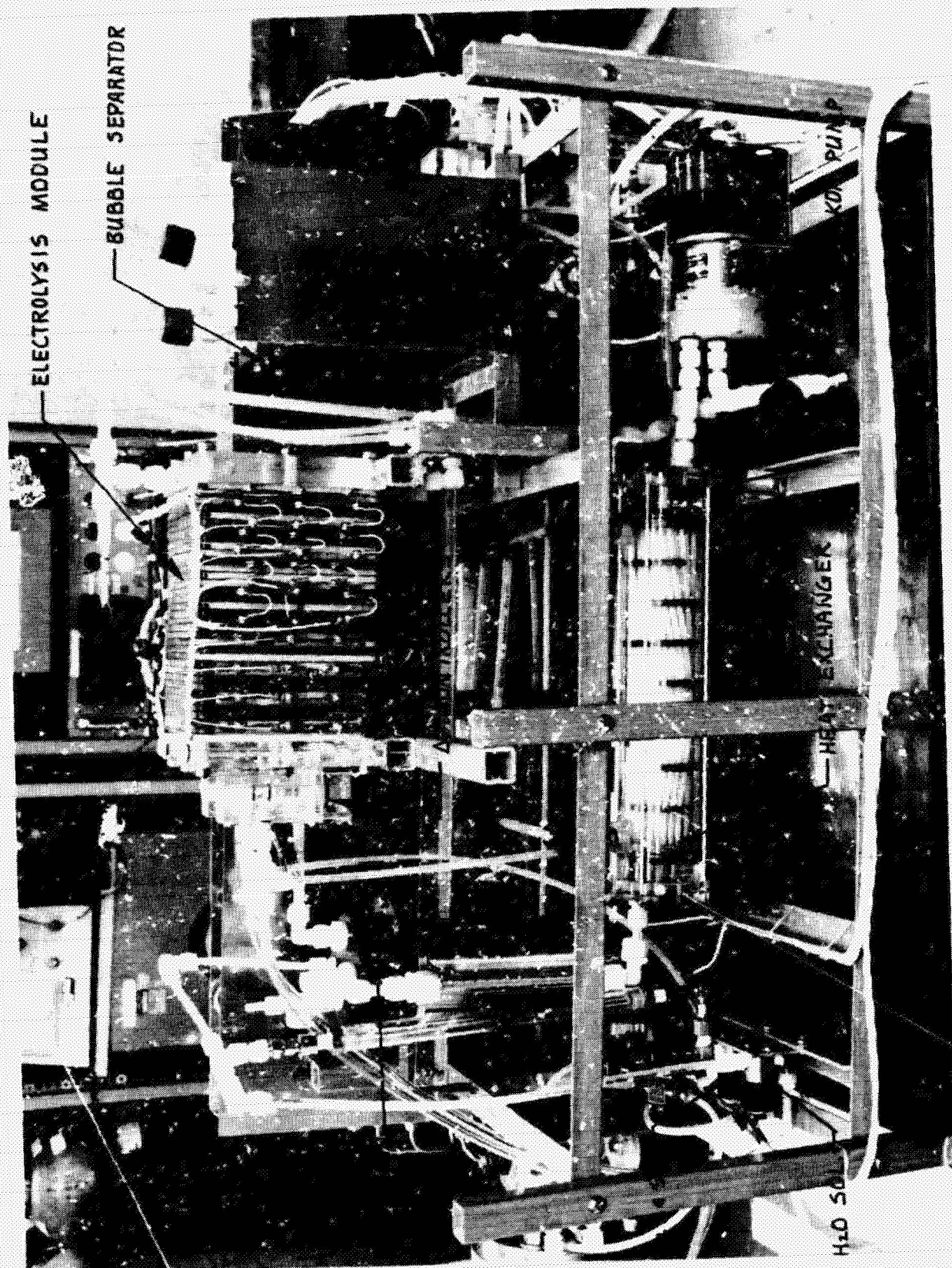


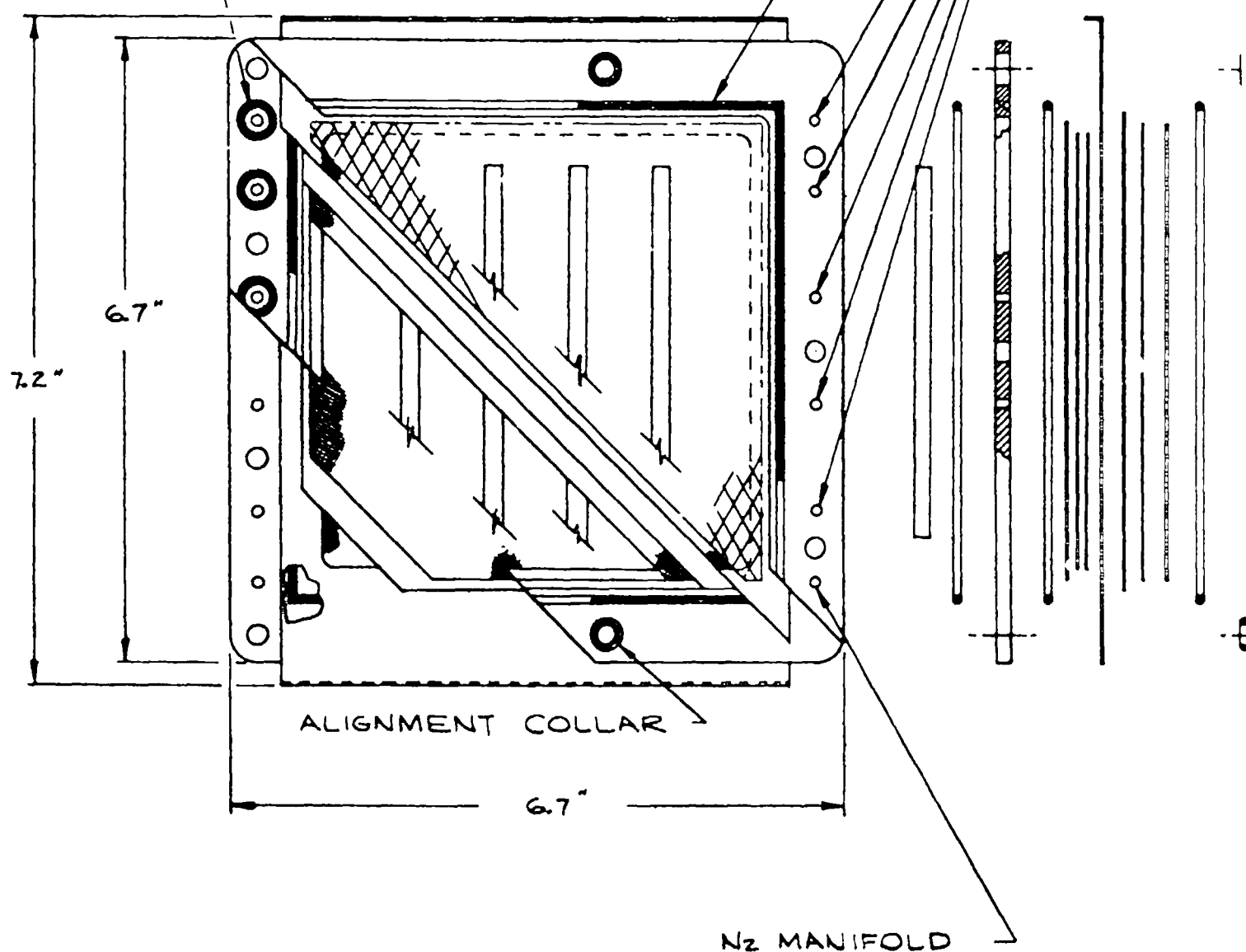
Fig. 2-16 O<sub>2</sub>/N<sub>2</sub> Generation System

O-RING FACE SEAL, NEAR AND FAR SIDE OF ELECTRODE SPACER, AROUND EACH MANIFOLD PORT.

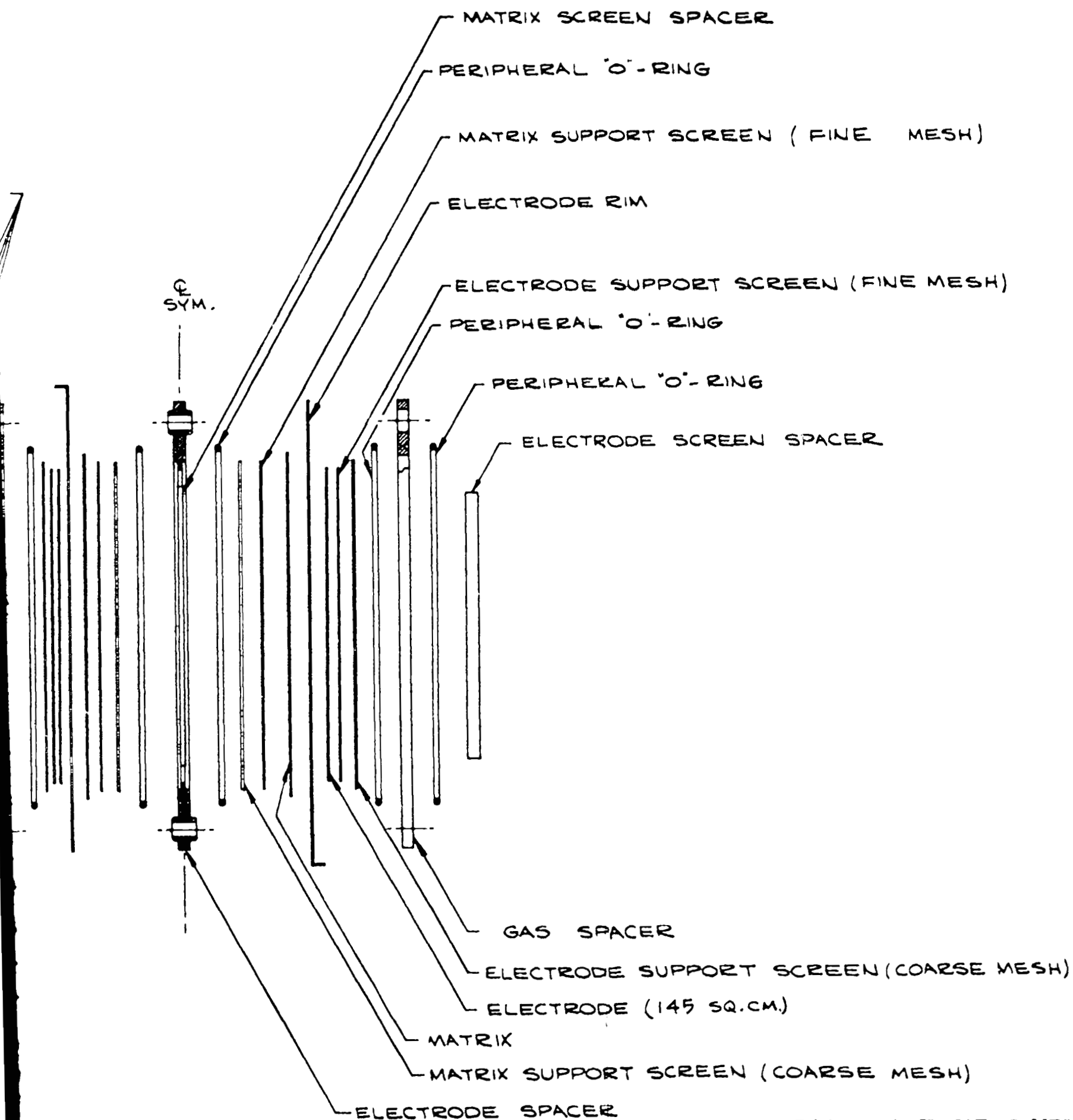
ELECTROLYTE MANIFOLD

N<sub>2</sub>/O<sub>2</sub> MANIFOLD

O-RING FACE SEAL, NEAR AND FAR SIDE, AROUND PERIPHERY OF ELECTRODE SPACER AND GAS SPACERS.







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Fig. 2-17 Cell Configuration

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2-33

FOLDOUT FRAME 2

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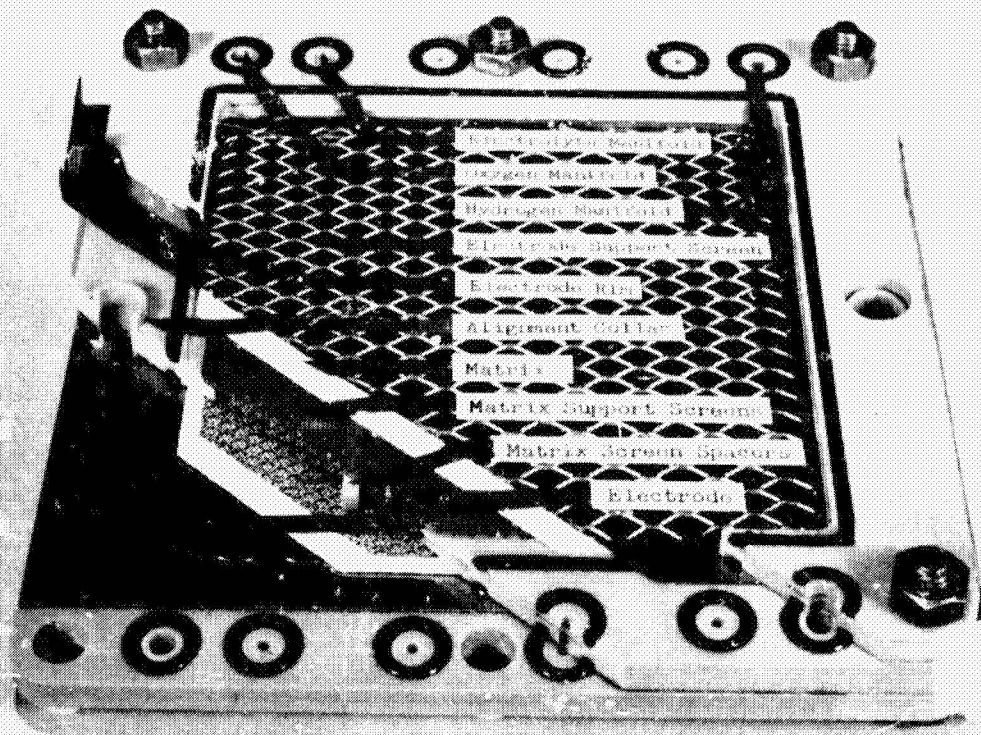


Fig. 2-18 Electrolysis Cell Cutaway View

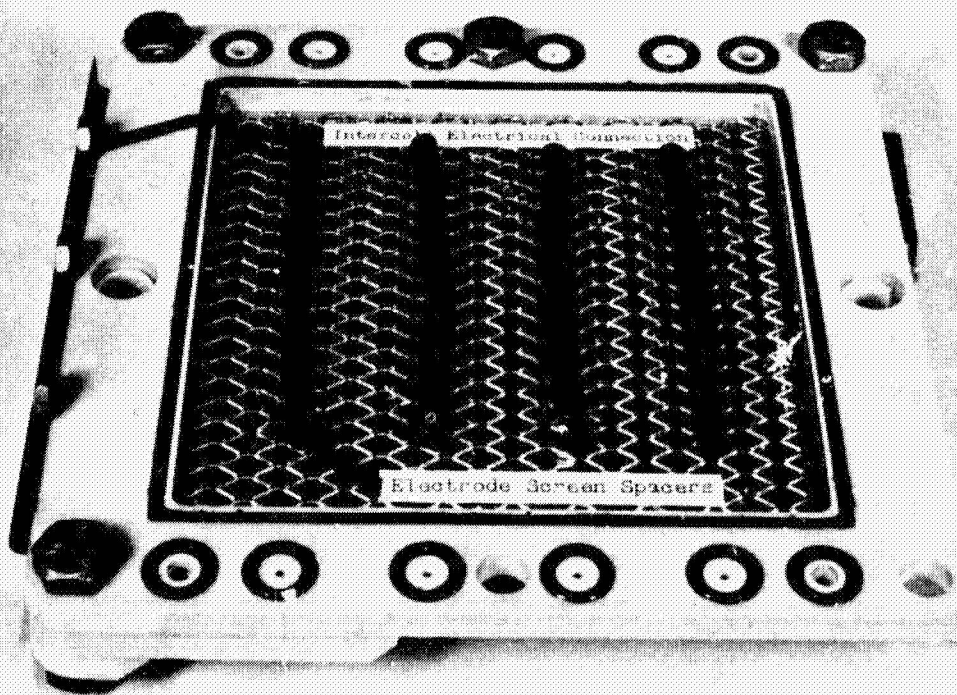


Fig. 2-19 Electrolysis Cell Reverse Side View

The matrix material is a highly refined asbestos. It is held in place between the electrode structure and a set of nickel support screens. Epoxy spacers located in the electrolyte space hold the opposing matrix support screens apart. Resilient elastomer material is used for pressure strips on the gas side of the electrode to ensure uniform contact between the matrix and the electrode. These strips also act as springs in the stack of cells to take up tolerances in the assembly.

The cells are arranged in the stack in a cathode-anode-anode arrangement so that each internal gas spacer serves two adjacent cells. This reduces the cell spacer components required per cell (except for one end cell) to one electrolyte and one gas spacer. When the stack of cells is being assembled, alignment collars on the electrolyte spacers key the cell parts in the proper position and prevent misalignment.

The 17 cells in the stack are connected hydraulically in parallel and electrically in two series banks of eight and nine cells.

Heat generated by the electrolysis reaction is removed from the cell with the circulating electrolyte. By removing the heat in the external loop of the circulation system, close control of cell temperature can be maintained. This permits control of the dew-points of the effluent gases below ambient, and thereby eliminates the problem of water condensation in gas lines, reservoirs, and valves downstream of the cell.

Water and  $N_2H_4$  consumed in the electrolysis reaction in the cell are replaced by direct injection of the proper liquid in the external electrolyte circulation loop. This method of makeup is conducive to rapid electrolyte equilibration even at high operating currents.

Differential Pressure Controllers. Two differential pressure controllers are mounted directly on one end plate of the cell stack. These devices sense the electrolyte pressure in the cell stack and throttle the hydrogen and oxygen/nitrogen effluent gases to provide 25 in. of water differential pressure, gas-over-liquid. When gases are not being generated in the cell stack, nitrogen purge solenoid valves open to provide gas



pressure. This differential pressure control is required to maintain the gas/liquid interface at the electrodes. The differential pressure controller assembly is shown in Fig. 2-20.

Bubble Separator. A bubble separator is located downstream of the cell stack in the electrolyte circulation loop, as shown in Fig. 2-15. The function of this device is to remove gas from the circulating electrolyte. The source of this gas is primarily dissolved gas in the feed water which is freed in the gaseous phase as water is consumed in the electrolysis reaction.

The bubble separator consists of hydrophobic and hydrophilic membranes assembled in a configuration, shown in Fig. 2-21, in which liquid can flow only through the hydrophilic membrane, and gas can flow only through the hydrophobic membrane. The gas side of the separator is vented through a differential pressure controller. This  $\Delta P$  controller is reverse-acting to the cell stack  $\Delta P$  controllers and is used to maintain the liquid pressure across the hydrophobic membrane higher than the gas pressure.

Electrolyte Pump. A pump is used to circulate the electrolyte. It is a commercial laboratory model consisting of a centrifugal impeller assembly which is driven with a magnetic coupling. All surfaces in the pump which contact the electrolyte are plastic.

Heat Exchanger. Waste heat generated in the cell stack is removed from the electrolyte, external to the cell stack, in a heat exchanger. The heat exchanger is a laboratory model shell-and-tube type device constructed entirely of plastic. Coolant is admitted to the shell side and electrolyte flows through the tube side.

Closed Reservoir. The reservoir is a variable-volume device designed for zero-gravity operation which is used to maintain system pressure. The configuration of this device is shown in Fig. 2-22. A rolling diaphragm works against a spring-loaded piston to maintain the pressure in the electrolyte circulation loop. Volume changes due to the water and hydrazine feed cycles are thus absorbed in the reservoir chamber.

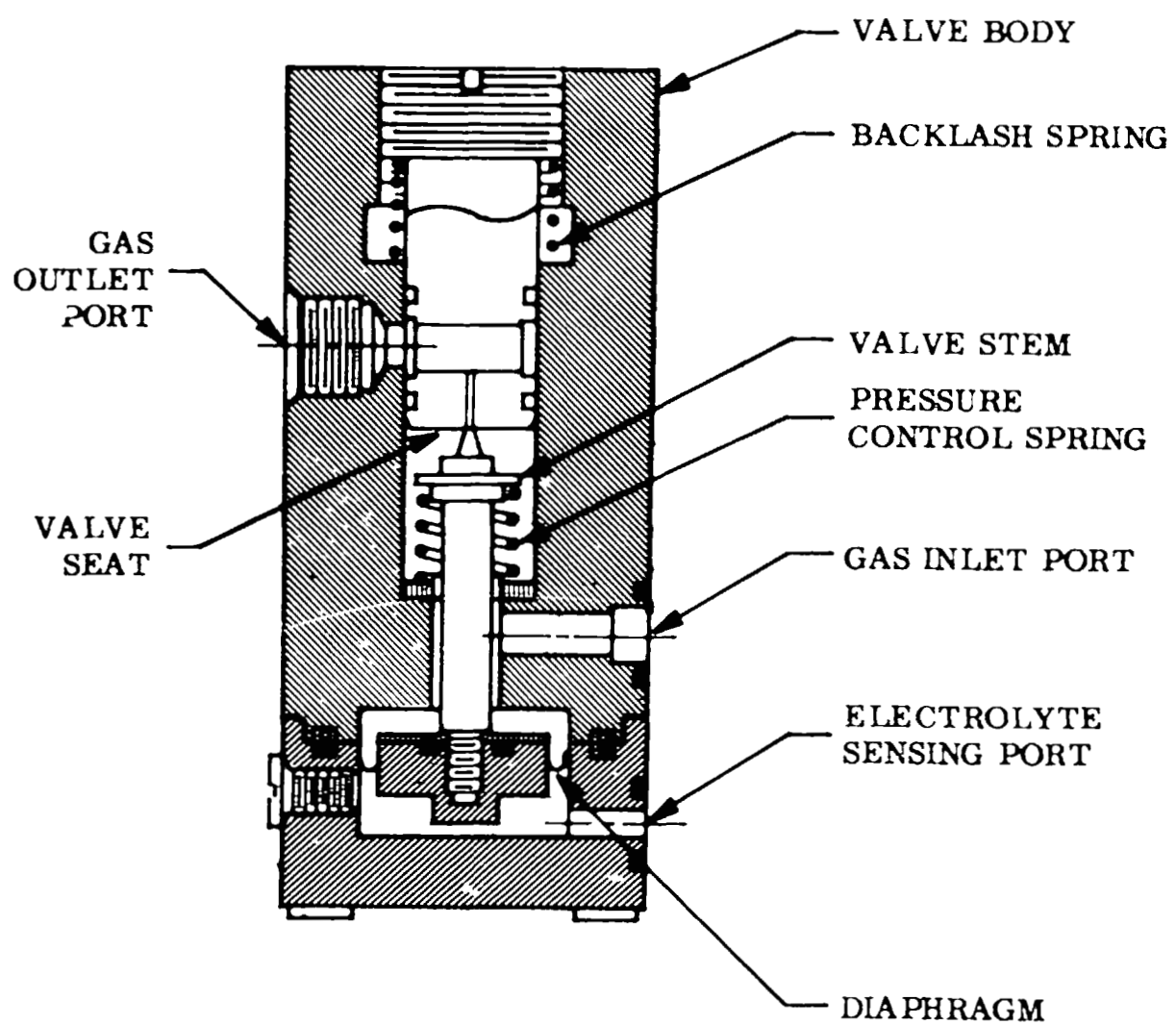


Fig. 2-20 Differential Pressure Controller Assembly

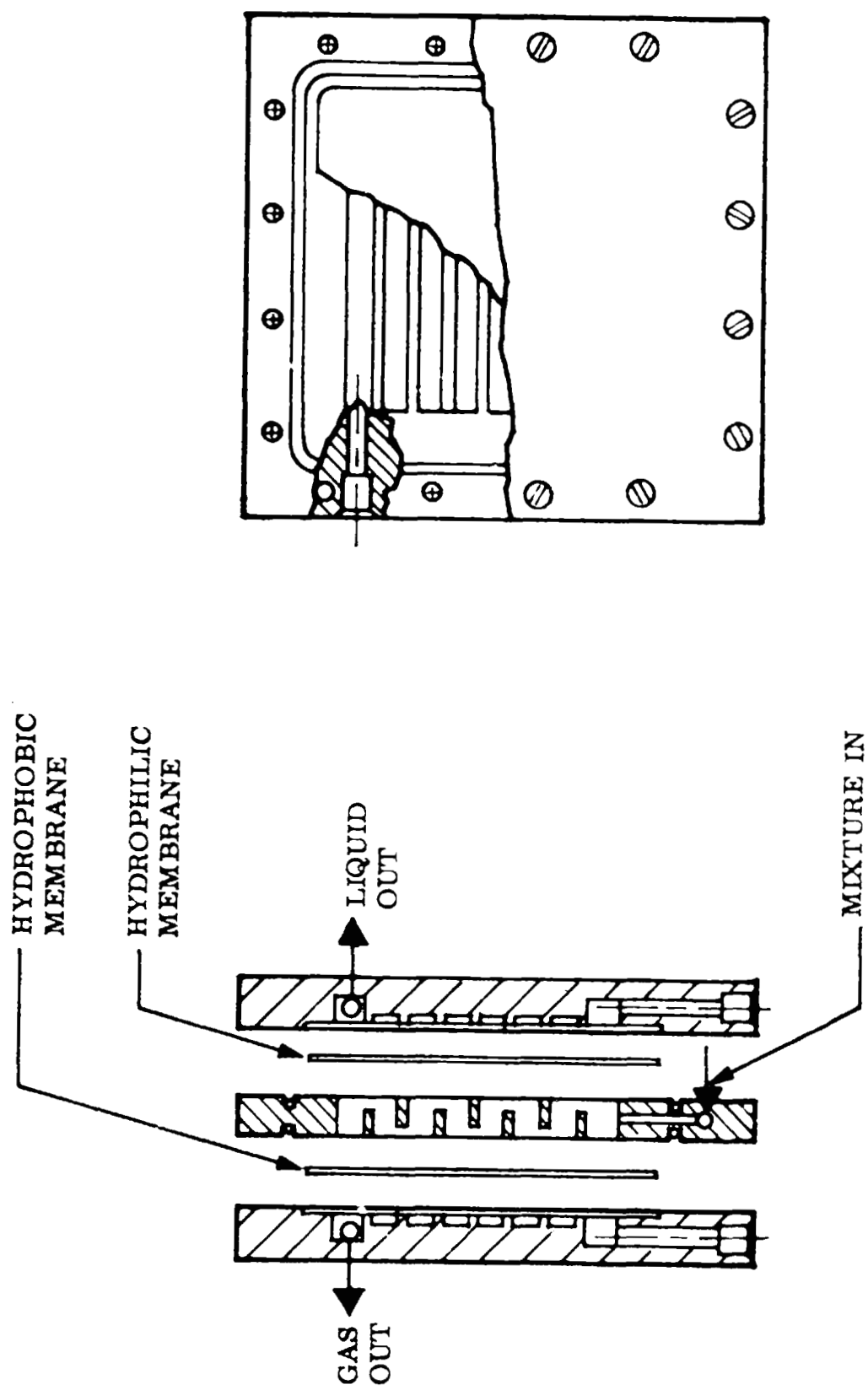


Fig. 2-21 Bubble Separator Configuration

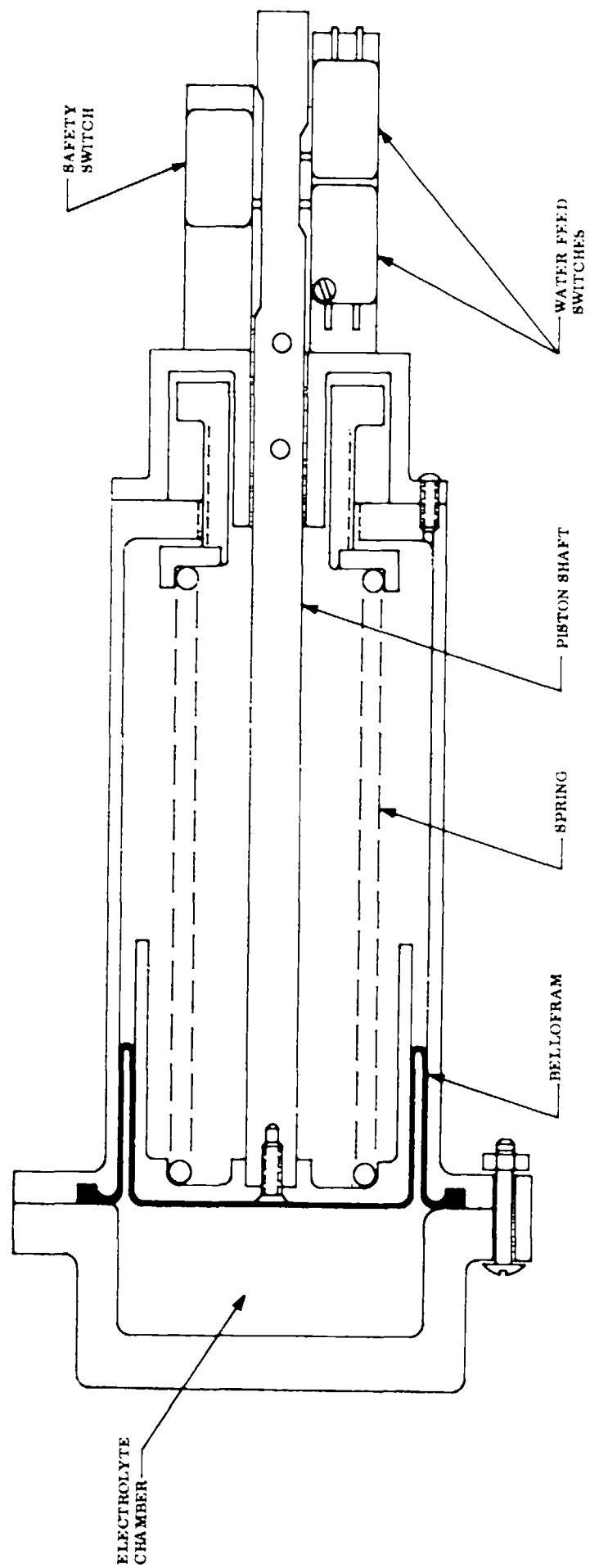


Fig. 2-22 Closed Reservoir Configuration

The position of the piston shaft is utilized to control the water feed; this control technique and the function of the safety switch are described in detail in Section 2.2.3.

Water and Hydrazine Storage. Spherical storage tanks for water and hydrazine are mounted underneath the oxygen/nitrogen generator. The water tank contains a silicone rubber bladder, and nitrogen gas pressure is applied to the back side of the bladder. When water feed is required, a solenoid valve opens in the water feed line, allowing the nitrogen pressure to force water into the closed reservoir. The hydrazine is stored in a similar tank from which the bladder was removed.\* It is mounted so that nitrogen pressure is applied at the top and hydrazine is withdrawn at the bottom of the tank. As with the water feed, a solenoid valve opens to allow hydrazine to enter the closed reservoir. A micrometer valve and flowmeter are provided to set the hydrazine flow rate. The plumbing required to permit filling, draining, and pressurization of these tanks is shown in Fig. 2-15.

#### 2.2.2 Cabin and Metabolic/Leak Simulator

The tank shown in Fig. 2-23 is used as a cabin simulator. It is equipped with gasket-sealed ports, an O-ring sealed door, and has a rotary fan mounted inside to assure gas-mixing. It was designed to be suitable for both vacuum and pressure applications. The volume of the tank is 30 cu ft.

The cabin simulator is integrated with the oxygen/nitrogen generation system and a metabolic/leak simulator system. It has, as well, an external sampling loop through which cabin gas is continuously circulated.

Metabolic/Leak Simulation. Metabolic consumption of oxygen and cabin leakage of oxygen and nitrogen are simulated by withdrawing gas from the cabin simulator at a controlled rate. Because oxygen cannot be withdrawn preferentially from the cabin,

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\*The bladder material, silicone rubber, is not compatible with hydrazine. A suitable ethylene propylene, required a special order, and delivery time was not compatible with the program schedule.

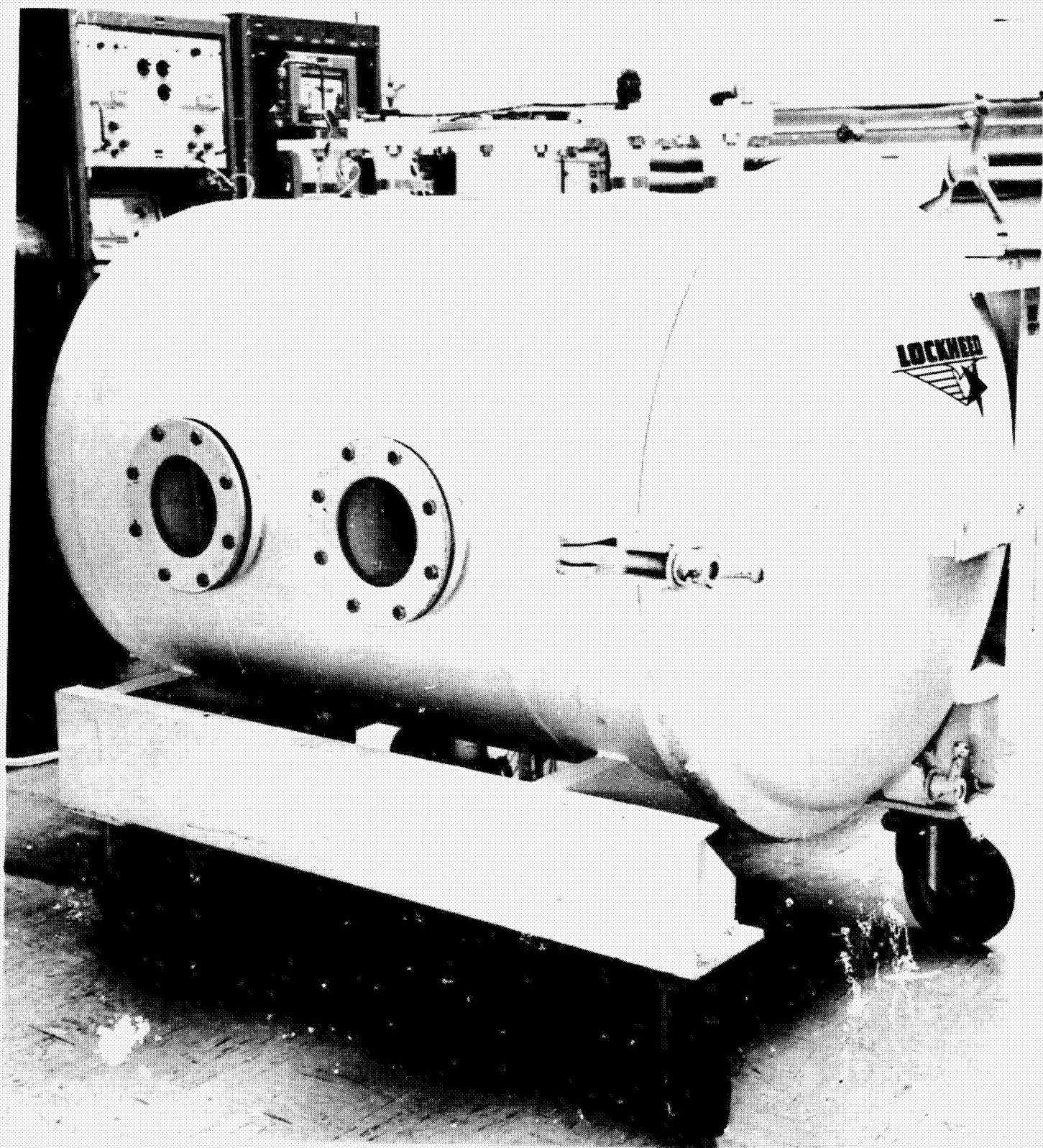


Fig. 2-23 Cabin Simulator

the total amount of gas removed includes the metabolic and leakage oxygen, the leakage nitrogen, and an amount of excess nitrogen removed with the metabolic oxygen. A controlled amount of nitrogen is replaced in the cabin simulator as makeup for excess nitrogen removed with the metabolic oxygen. The flow control valve and flow-meter configuration used to accomplish this metabolic/leak simulation is shown in Fig. 2-15.

Cabin Atmosphere Sampling Loop. The cabin simulator is operated below ambient pressure at approximately 12 psia. In order to remove the metabolic/leak gas, a diaphragm pump is used to raise the pressure in a sampling loop outside the cabin to a pressure above ambient. A nonrelieving pressure regulator is used to control the pressure in this loop. Cabin gas is continuously circulated through the loop for gas composition monitoring. Provision also is made for introducing calibration gases in the portion of the sampling loop containing the gas-monitoring sensors.

### 2.2.3 Instrumentation

The model system is instrumented with automatic controls of the oxygen-nitrogen generator functions, cabin atmosphere control, power conditioning and both digital and analog readouts of system parameters.

All control logic and power conditioning are handled with solid-state circuitry. The system is also equipped with automatic safety circuits which will cause complete system shutdown if unsafe conditions occur. A block diagram of the system instrumentation is shown in Fig. 2-24. Detailed circuit diagrams are included in Appendix A.

#### 2.2.3.1 System Controls and Safety

Water balance in the electrolyte loop is maintained by controlling the electrolyte volume. A set of microswitches turn the water feed solenoid valve on and off as the piston in the closed reservoir moves up and down. The movement of the piston shaft and the actuation

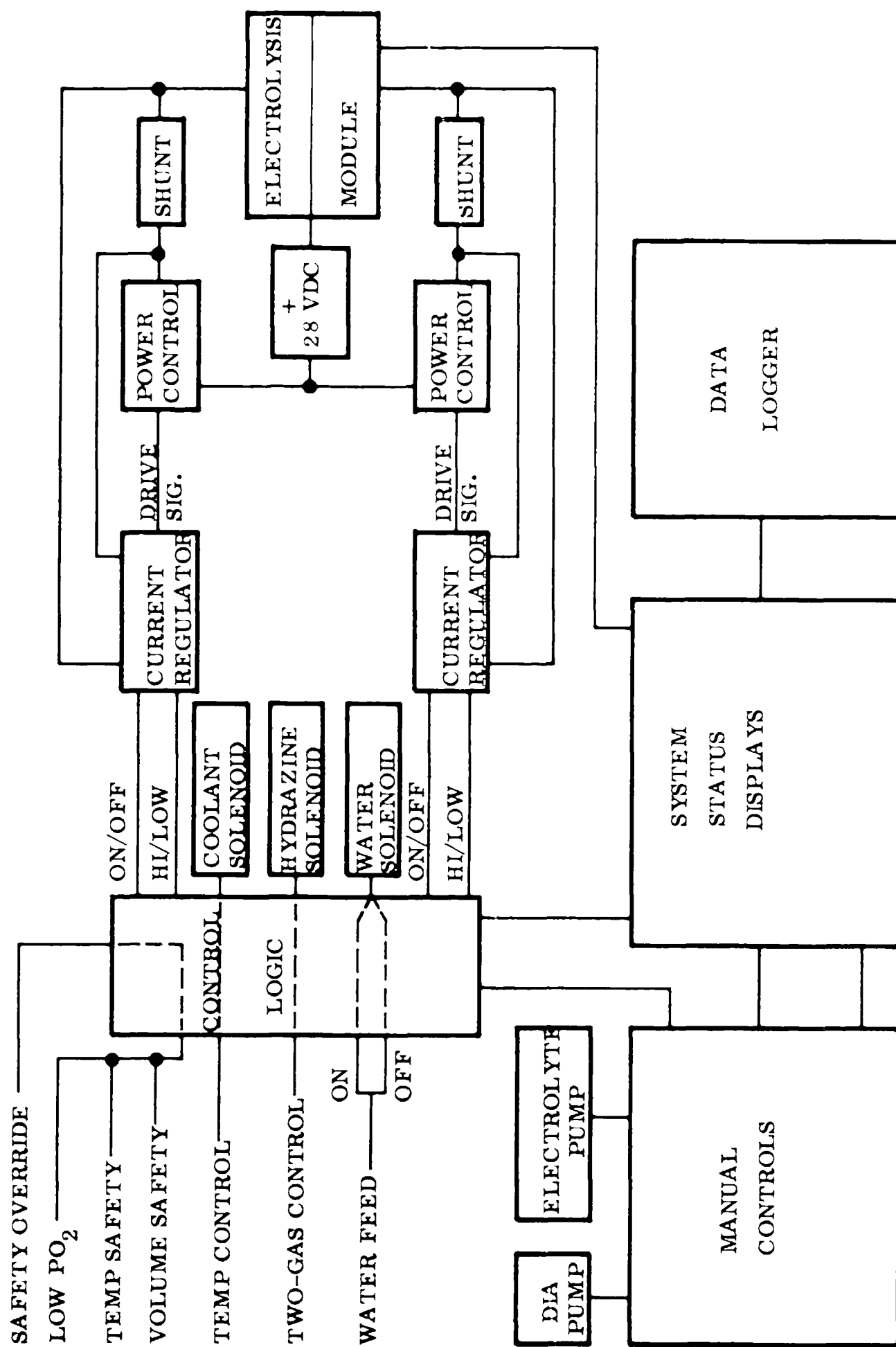


Fig. 2-24 O<sub>2</sub>/N<sub>2</sub> System Control Block Diagram



of the microswitches by a detent in the shaft that occurs during a water feed cycle is shown in Fig. 2-25. In position (a), the piston is in the middle of the water feed control band. As water is consumed in the electrolysis reaction, the electrolyte volume shrinks and the piston travels upward to position (b). Here the lower microswitch closes and signals the water feed solenoid to open. As water feed continues, the piston travels down through position (c) and continues to position (d), where closure of the upper microswitch signals water feed shutoff. The cycle described in this figure is completed as the piston returns to its starting position at (e).

The piston travels approximately  $3/8$  in. during a water feed cycle with a corresponding volume change of about 45 cc.

The control logic is such that water feed is commanded off whenever both microswitches are actuated, as would be the case if the piston shaft detent is completely above or below both switches. A manual override is provided so that the operator can command the water feed either on or off at any time.

Control of the electrolyte temperature, necessary because of the waste heat generated in the electrolysis reaction, is accomplished by using a thermost switch in the electrolyte discharge from the cell stack to provide a control signal to a coolant solenoid valve. On demand, the solenoid valve opens to allow coolant (ethylene-glycol) to flow through the electrolyte heat exchanger. The flow rate is set with a flow control valve. Control of the electrolyte temperature also provides control of the dewpoints of the generated gases. The thermost switch used in this system has a switch-closure setting of  $75^{\circ}\text{F}$ .

On-off control is provided for the hydrazine feed. A polarographic oxygen partial pressure sensor in the cabin simulator sampling loop provides the signal to the hydrazine-feed solenoid valve. When cabin  $\text{PO}_2$  reaches an adjustable upper set point, the hydrazine solenoid opens to admit flow to the closed reservoir. The solenoid valve remains open until cabin  $\text{PO}_2$  reaches an adjustable lower set point. A manual micrometer valve and flowmeter can be used to set the hydrazine flow rate during the

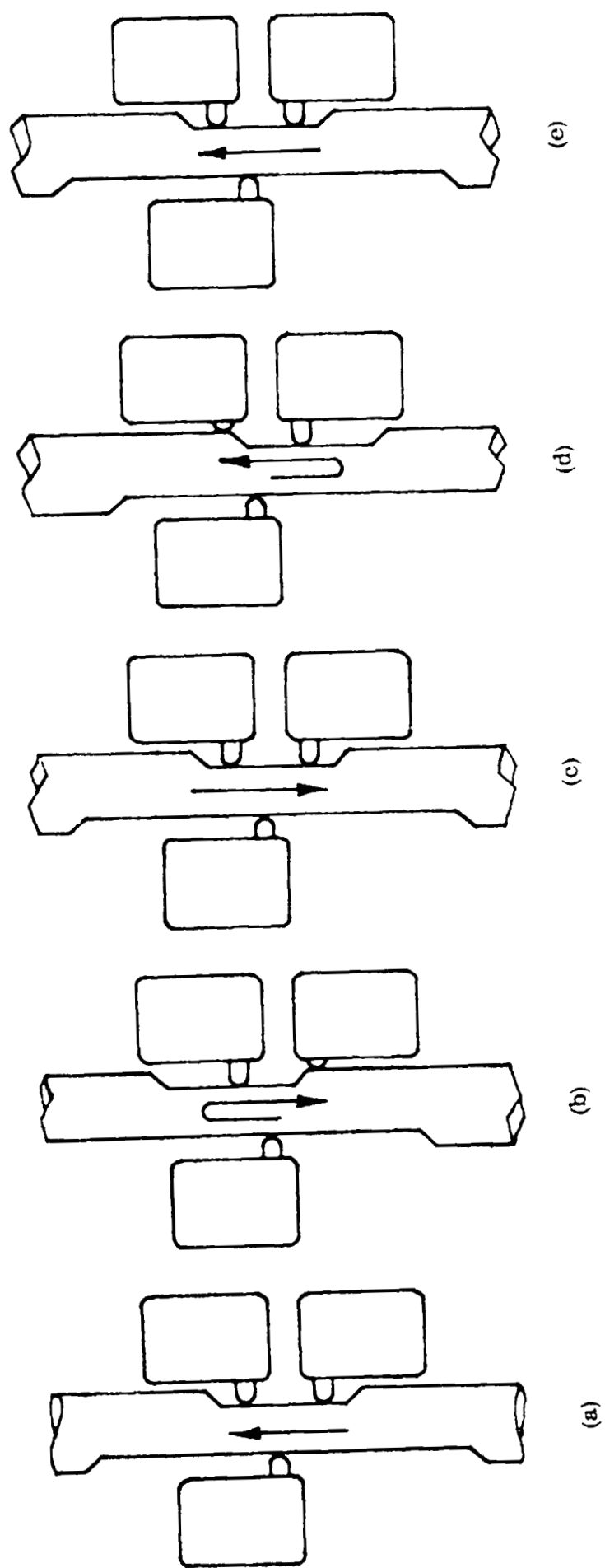


Fig. 2-25 Water Feed Control Mechanism

on portion of the feed cycle. Manual override is provided so that hydrazine can be commanded on or off at any time.

The electrolysis cell stack current control utilizes a high/low mode of operation. Both the high and low current values are adjustable. A signal from a pressure transducer is used to control the current mode. When the cabin total pressure ( $P_T$ ) reaches an adjustable upper limit, the current is commanded to the low mode. The low mode is maintained until  $P_T$  reaches an adjustable lower set point; the current then switches back to the high mode. Manual override is provided so that the current mode can be commanded low, high, or off at any time.

In order to minimize the operating temperature of solenoid valves used in this system, a holding-current technique is employed. Whenever a signal to energize a solenoid valve coil occurs, a brief actuation current pulse is provided and then the current is reduced to a holding level. Power dissipation with this holding current is sufficiently low so that the solenoid valve bodies remain essentially at ambient temperature. This approach is especially useful for solenoid valves that remain energized closed for long periods, and increases the operating life of the valve seals.

Sensors and circuit logic are provided to effect automatic shutdown of the system in the event of an unsafe or abnormal operating condition. The shutdown logic, when actuated by a safety sensor signal, disables the electrolyte pump, turns off the water and hydrazine feed, opens the  $N_2$  purge solenoid valves to purge the  $H_2$  and  $O_2/N_2$  chambers of the cell stack, and diverts the  $O_2/N_2$  module effluent from the cabin simulator to a fume hood.

One of the safety sensors is located on the closed reservoir. Recalling that the water feed control utilizes a set of microswitches which operate on a detent on one side of the reservoir piston shaft, a similar detent and associated microswitch on the opposite side of the piston shaft provide a high or low reservoir volume signal. The safety switch is set to actuate  $\pm 150$  cc on either side of the water feed control band. High-volume shutdown would occur in the event of overfeed of water or hydrazine. Low-

2

volume shutdown would occur in the event of lack of water feed or a leak in the electrolyte loop.

Overtemperature protection is provided in the form of a thermostatic switch mounted in one end plate of the cell stack in contact with an end electrode. This sensor is a two-position sensor that actuates a warning light at 85°F and an automatic system shutdown at 105°F. Overtemperature shutdown would occur in the event of coolant supply failure or electrolyte pump failure.

Protection against excessive hydrazine feed is provided by a polarographic  $\text{PO}_2$  sensor in the cell stack  $\text{O}_2/\text{N}_2$  effluent gas line. When an adjustable minimum effluent  $\text{PO}_2$  set point is reached, the hydrazine feed solenoid valve is automatically shut off.

Power loss protection is provided. In the event of the failure or even momentary loss of plant power, the system will automatically shut down and will not restart after power has been restored until manually reset.

The system control and safety instrument panel is shown in Fig. 2-26. The lefthand side of the panel contains all of the electronic instrumentation; the center panel has the data logging signal leads; and the righthand side contains the flow controls for the cabin and metabolic/leak simulation.

Fault diagnosis is provided by the upper row of lights on the instrument panel, which can be seen in this figure. Indicators include temperature warning, overtemperature shutdown, high/low volume shutdown, and safety  $\text{PO}_2$  shutoff of the hydrazine feed. The safety indicator that indicates the cause of automatic shutdown will remain on until the system is manually reset. All of the safety circuits contain logic to prevent the system from being reactivated automatically before the cause of the unsafe condition can be rectified.

A manual override of the safety circuits is provided and is used during system startup at the beginning of a test.

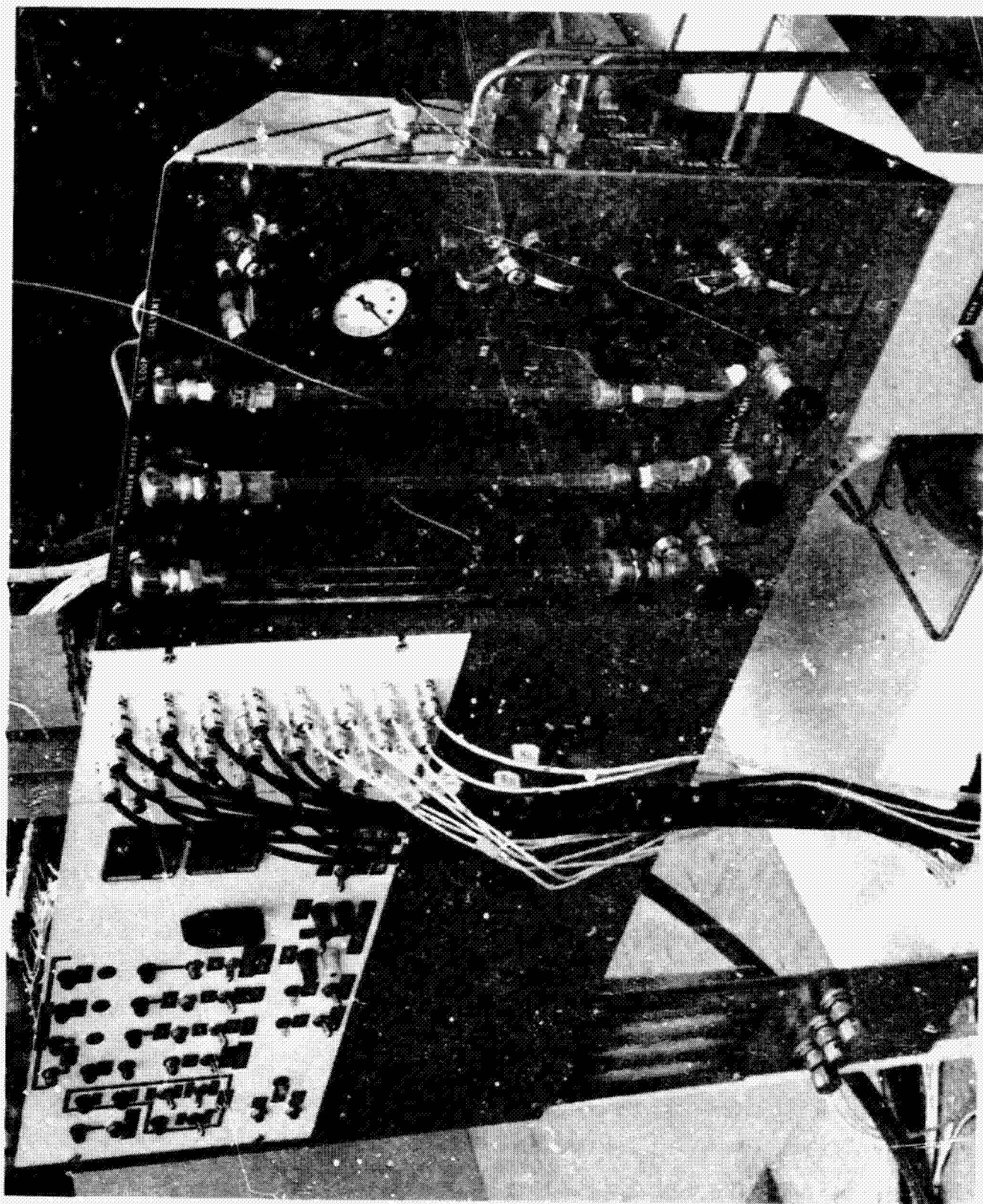


Fig. 2-26 O<sub>2</sub>/N<sub>2</sub> System Control Panel

#### 2.2.3.2 Power Conditioning

Conditioning and regulation of the power for the electrolysis cell stack is accomplished with high-efficiency programmable current-switching regulators. A pulsewidth-modulated signal drives a power-switching amplifier, which produces a constant current through the cell stack.

The 17-cell stack is divided into an eight- and nine-cell electrical bank. Each bank has its own current regulator, but shares outputs from the current mode controller and oscillator.

Figure 2-27 illustrates the technique employed to obtain this type of regulation. Load current is sampled via a differential voltage developed across the shunt resistor,  $R_S$ . This signal is amplified by the feedback amplifier and presented to the error amplifier where it is compared with a fixed reference signal. If a difference exists between the sampled signal and the reference signal, an error voltage will be developed and fed to the pulsewidth modulator. A fixed frequency (approximately 15 kHz) squarewave oscillator provides the trigger source for the pulsewidth modulator. The output of the modulator drives the switching transistor,  $Q$ .

The transistor serves as a switch and is either shut off or saturated. The length of time the transistor is "on" or saturated depends on the time length of the pulse emitted from the modulator. The operation of the circuit is divided into two cycles: first, when transistor  $Q$  is on; second, when  $Q$  is off.

During the interval when  $Q$  is on, a current path is provided from the positive side of the unregulated input source through inductor,  $L$ , shunt,  $R_S$ , the cell bank load, and through the transistor to the supply return. Switching diode,  $D$ , is back-biased during this interval; thus, no appreciable current flows through it. Capacitor,  $C$ , is allowed to charge to the cell bank voltage, thus storing energy. Since the voltage across the nonsaturating choke inductor,  $L$ , is constant, the current through the

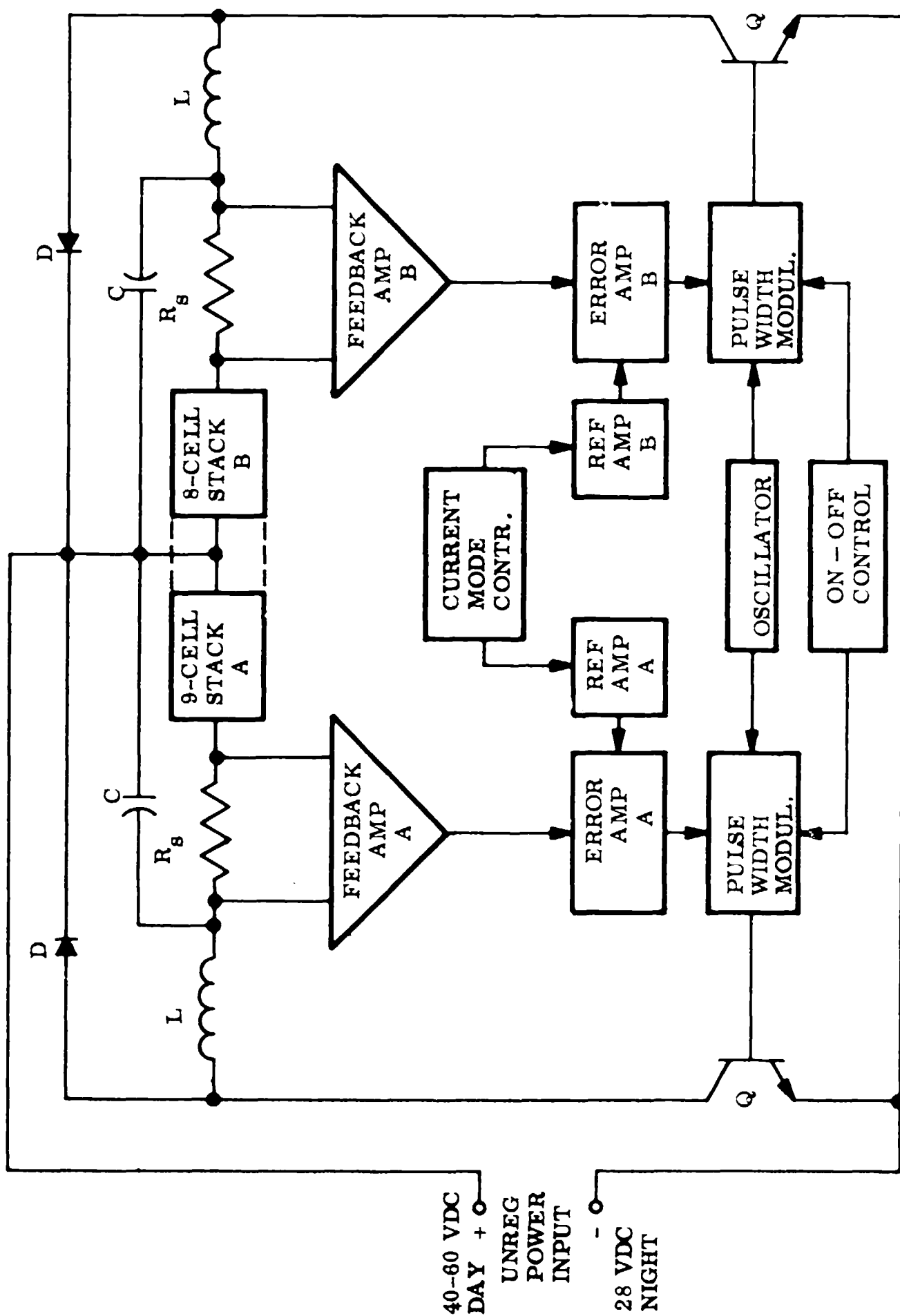


Fig. 2-27 Power Conditioning Technique



choke - which is also the shunt, load, and transistor current - will increase linearly with time according to  $e_L = L di/dt$ . When the average current through the shunt equals the reference signal current level, the error is decreased to zero and the modulator turns the switching transistor off, which triggers the second cycle of operation.

When the current source through the choke is inhibited by the transistor switch, the voltage across the choke immediately reverses due to the decreasing choke current, thus forward-biasing and turning on switching diode, D. At this point, the choke and capacitor start to release their stored energies, thus providing a current source through the diode to the cell-bank load for this half of the cycle. When the average load current falls below the reference current level, the transistor is switched on again via the pulsewidth-modulator/error amplifier, and the cycle repeats.

The load currents are programmed by varying the reference current signal level to obtain high- and low-mode currents.

The efficiency of this circuit is inherently high as compared with conventional series regulators because the power losses are minimized by saturating the switching transistors and diode when they are turned on and not allowing the choke to become saturated.

Current regulation components of this circuit are mounted on a circuit card which plugs in behind the control panel. High-current components are mounted on an air-cooled cold plate, which can be seen in Fig. 2-28.

#### 2.2.3.3 Control and Monitoring Instrumentation

The model system is instrumented with the sensors required to provide signal inputs for the control, safety, and power conditioning circuits described above. In addition, sensors and readouts are utilized to provide system performance data. The sensing and readout techniques used in the system are given in Table 2-2. Some of the instrumentation is shown in Fig. 2-29.



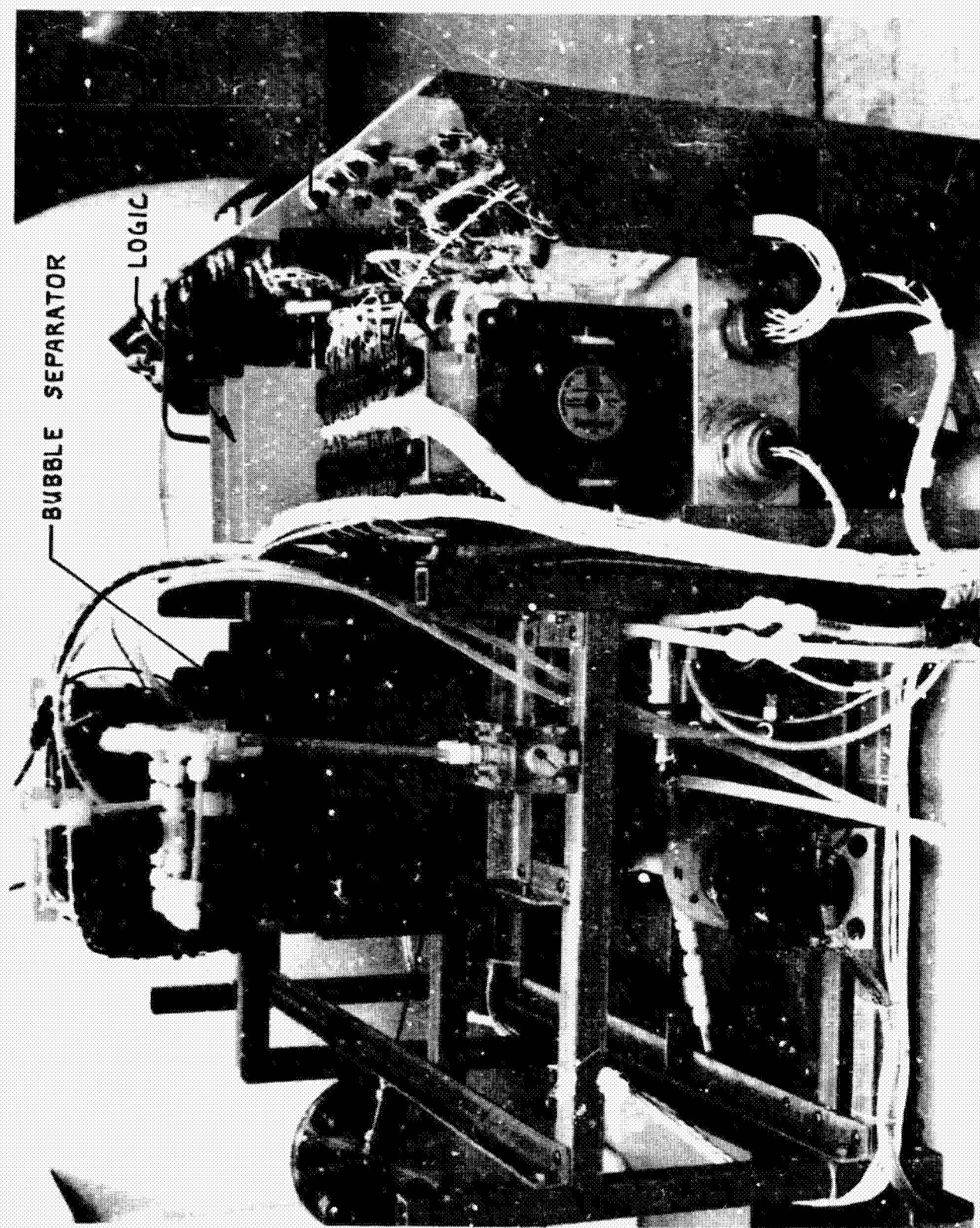


Fig. 2-28 O<sub>2</sub>/N<sub>2</sub> System Side View

Table 2-2  
SYSTEM CONTROL AND MONITORING INSTRUMENTATION

Parameter	Measurement Technique	Signal Use
Cabin total pressure	Transducer Transducer Vacuum gage	Current mode control Strip-chart recorder Transducer check
Cabin oxygen partial pressure	Polarographic analyzer Polarographic analyzer Paramagnetic analyzer	Hydrazine feed control Strip-chart recorder Polarographic sensor check
Effluent oxygen partial pressure	Polarographic analyzer	Strip-chart recorder and PO <sub>2</sub> safety
Hydrazine feed valve on-off status	Electrical impulse	Strip-chart recorder
Electrolyte temperature	Thermoswitch	Coolant solenoid valve control
Module temperature	Thermoswitch 2-position	Overtemperature warning and shutdown
Coolant temperature	Thermometer	Data logging
Cell bank current	Shunt and digital voltmeter	Data logging
Cell bank voltage	Digital voltmeter	Data logging
Individual cell voltages	Digital voltmeter	Data logging
High/low current mode time	High/low mode timers	Data logging
Metabolic/leak flow rate	Rotameter and wet test meter	Data logging
Nitrogen makeup flow rate	Rotameter	Data logging
Hydrazine flow rate	Rotameter	Data logging
Electrolyte circulation rate	Rotameter	Data logging
Coolant flow rate	Rotameter	Data logging
Hydrogen production	Wet test meter	Data logging
Electrolyte volume	Microswitch - piston position Microswitch - piston position	Water feel control High/low volume safety
Cabin sampling loop pressure	Pressure gage	PO <sub>2</sub> sensor calibration
Module pressure	Pressure gage	Data logging
Hydrazine tank pressure	Pressure gage	Safety check
Bubble separator gas evolution rate	Water volume displacement	Data logging
Input electrolysis power	Power supply voltmeter and ammeter	Data logging

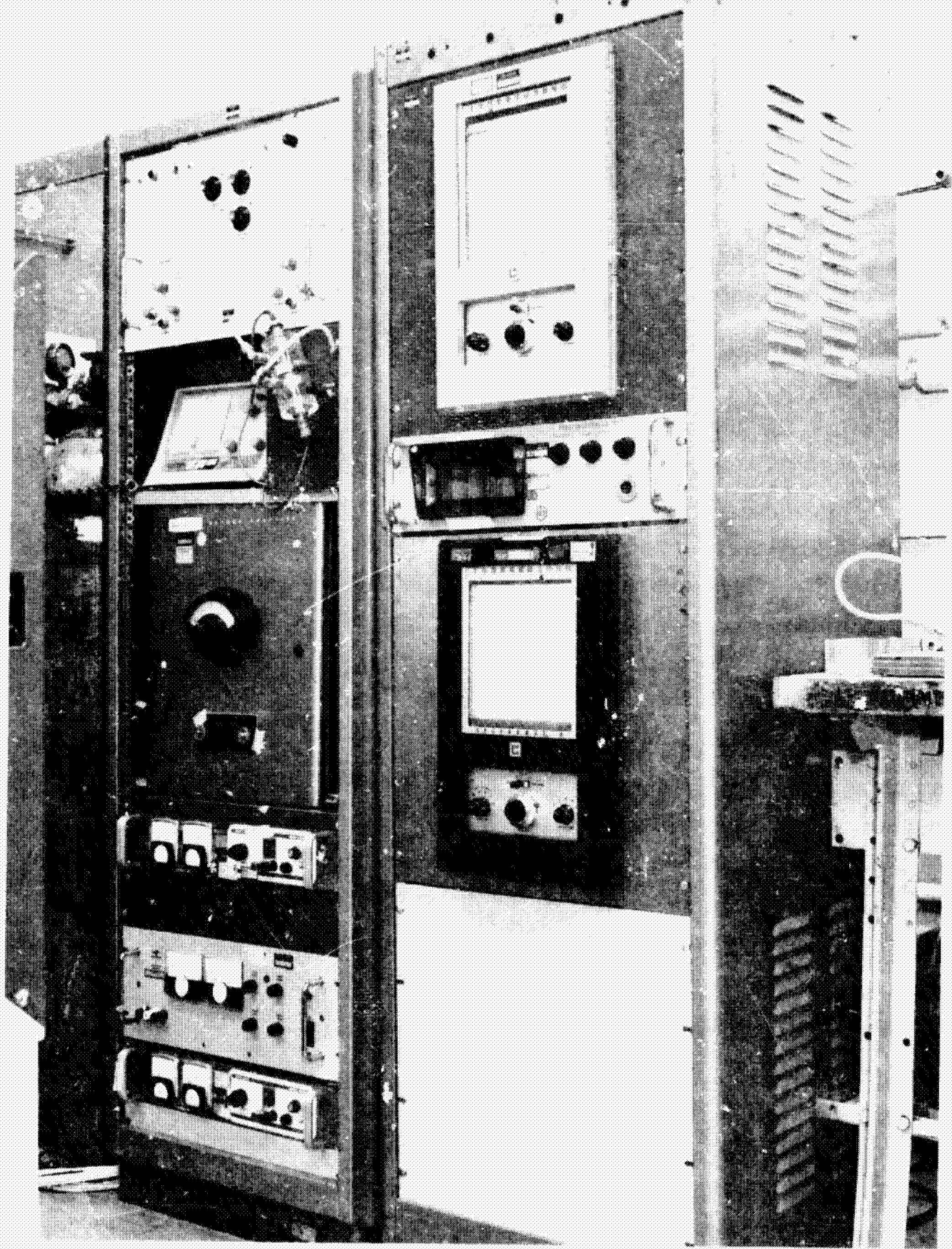


Fig. 2-29 Test Instrumentation



## 2.3 SYSTEM TESTING

Testing of the one-man model  $O_2/N_2$  system was conducted under this program with the objectives of evaluating the performance of various system components, establishing the characteristics of the cabin atmosphere control that can be achieved with this technique, and providing experimental data for the updating and verification of a computer routine model of the system.

### 2.3.1 Operating Procedure

The test plan that was established for meeting the objectives stated above involved a series of tests. Each test was conducted in a continuous operating mode with around-the-clock monitoring. The duration of each test was determined by the specific data requirements. Long-duration testing of the basic electrolysis system in previous programs (Ref. 3) has already established the capability of a system of this type to operate continuously for periods exceeding one year.

Startup-Shutdown Procedures. The detailed procedure for starting up and shutting down the system is described in Appendix B. The general approach to startup was to establish control conditions as rapidly as possible so as to allow the major part of a week for operation under automatic control conditions. This required that the cabin simulator be evacuated by increasing the metabolic/leak flow and manually adjusting the  $N_2$  makeup flow to maintain the cabin  $PO_2$  near the control band. At the same time, the hydrazine feed rate had to be increased to enrich the hydrazine concentration in the electrolyte.

Data Logging. While only the hydrazine concentration in the electrolyte, cabin total pressure, and cabin oxygen partial pressure were required data for computer routine analysis, additional module and system status data were recorded on an hourly basis to allow a thorough analysis of the system operating characteristics. The test data logs are included in Appendix C.

Test Monitoring. Around-the-clock monitoring was provided during system testing. The metabolic/leak and nitrogen makeup flow rates, which tended to drift, were adjusted as necessary. The cabin and effluent  $PO_2$  sensors were calibrated every 8 hr in the first test, but little drift was observed. In subsequent tests, these sensors were calibrated at the beginning and end of each test.

The water storage tank was filled with a known volume at the beginning of each test and the residual was measured at the end. Hydrazine consumption was measured by draining and back-filling the hydrazine tank at the end of each cabin  $PO_2$  cycle.

Gas and Electrolyte Sampling. Periodic sampling of the effluent gases was conducted to verify the  $PO_2$  sensor readings. The samples were analyzed using gas chromatographic techniques.

Electrolyte samples were taken just prior to the closing of the hydrazine solenoid valve, to obtain the maximum hydrazine concentration, and just prior to the opening of the feed valve, to obtain the minimum hydrazine concentration. Chemical analysis of the electrolyte samples for hydrazine content utilized the direct iodate method with solvent (Ref. 6).

### 2.3.2 System Test Summary

A series of four tests was conducted on the one-man model  $O_2/N_2$  system. The test sequence logic is shown in Fig. 2-30.

The objectives of the first test were to check out the operation of the fully integrated  $O_2/N_2$  generation system as well as the cabin and metabolic/leakage simulator, and to establish the test procedure for subsequent tests. Highlights of this test include:

- Successful operation of the zero-gravity closed reservoir and bubble separator.
- Verification of  $P_{total}$  control for a fixed metabolic and leak rate at a control band of 12.28 to 12.30 psia.

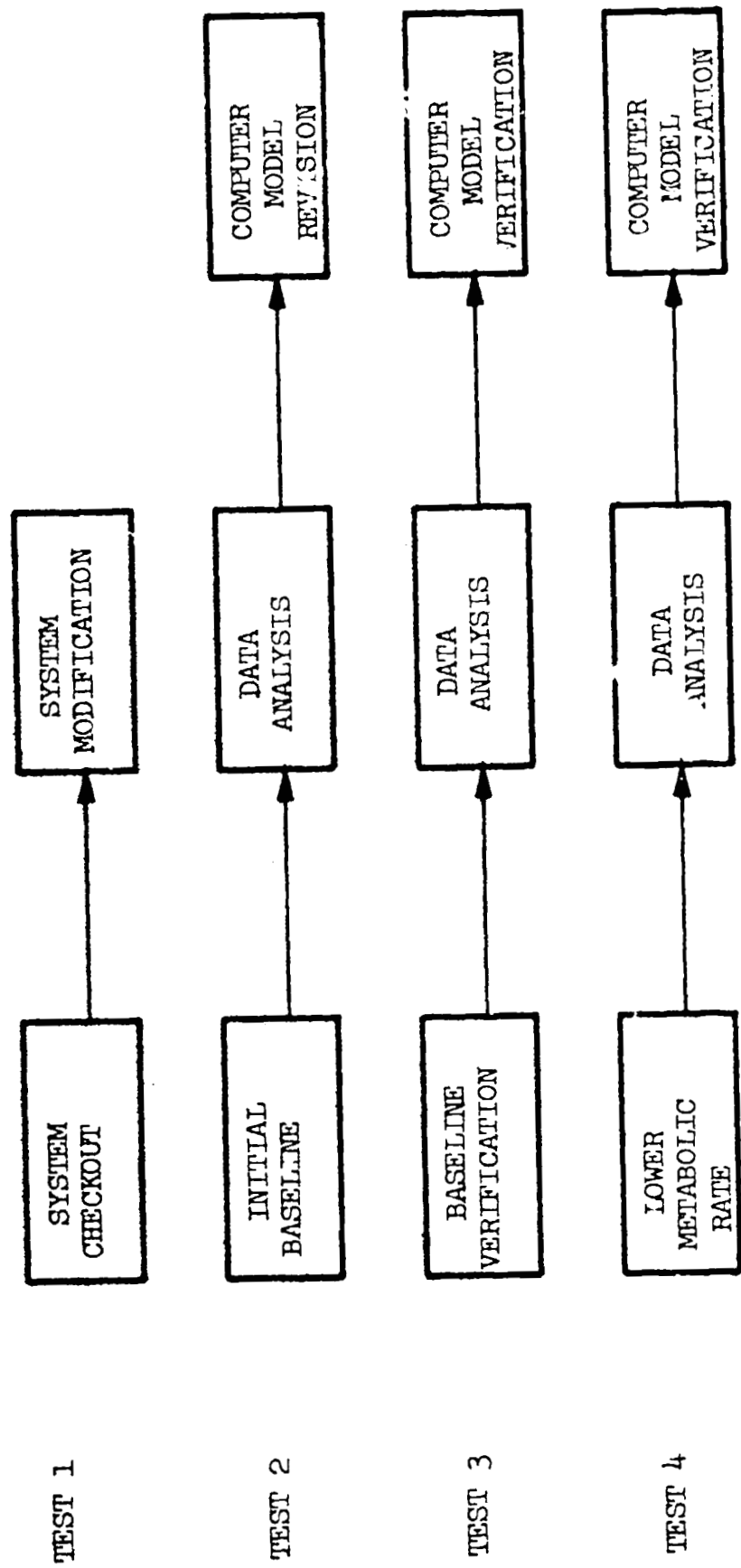


Fig. 2-30 System Test Sequence

- An indication of cabin  $PO_2$  control at a control band of 2.85 to 2.95 psia. A full  $PO_2$  cycle was not completed.
- Nitrogen in the effluent oxygen measured over a range from 0 to 91 percent.

Several system and procedure modifications indicated by this test were implemented before proceeding to the next test. These modifications are discussed in detail in Section 2.3.3.

Test 2 was the initial baseline run of the system at the design conditions; that is, at a one-man metabolic load with a one-man proportion of a space cabin leakage. Summary results of this test showing the cabin total pressure and oxygen partial pressure control are given in Fig. 2-31.

The baseline was rerun in Test 3 to provide data for verification of the computer model revision. The summary results of this test are shown in Fig. 2-32.

In Test 4, both the high/low currents and the metabolic load were varied to provide a different set of data points. In the one-man system, changes in metabolic load were made, approximately corresponding to reducing the crew size from 12 men to 3, 6, and then to 10 men. Summary results are shown in Fig. 2-33.

### 2.3.3 Test Results and Discussion

#### 2.3.3.1 Test Logs and Performance Data

Test 1 Results. An operational checkout test was conducted for a duration of 86 hr. The Time/Event Log for this test is presented in Table 2-3.

Cabin  $PO_2$  and hydrazine concentration for the period from 46 to 81 hr elapsed time are shown in Fig. 2-34. The shaded area in this figure is the control signal band that was shifted at 54 hr to correct for the higher pressure in the sensor canister.  $PO_2$  control capability was demonstrated by the turnaround of the cabin  $PO_2$  at approximately 61 hr. A full cycle of the  $PO_2$  was not achieved because of a problem with the water feed, which occurred at 81 hr.

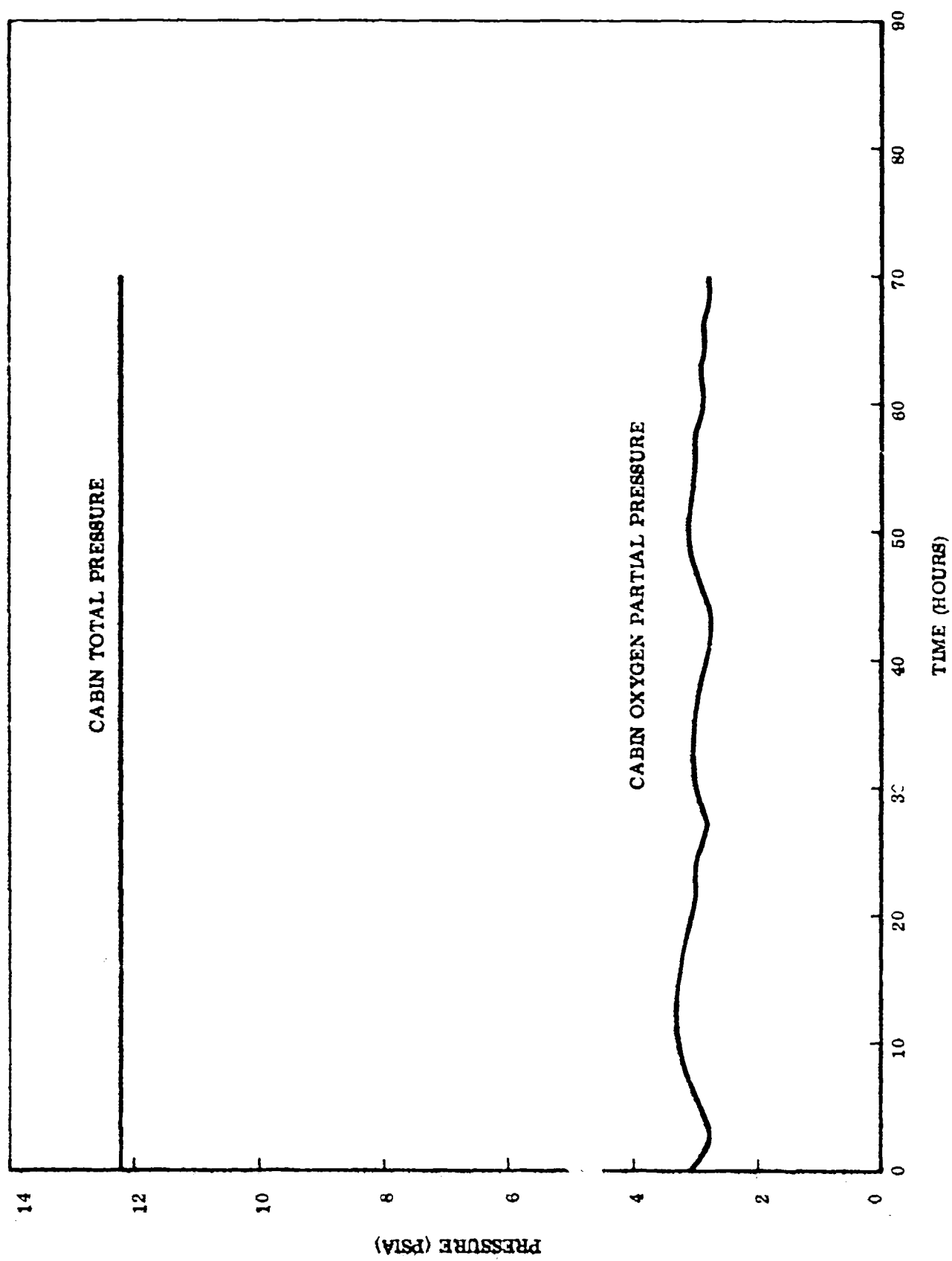


Fig. 2-31 Test 2 - Cabin Atmosphere Control



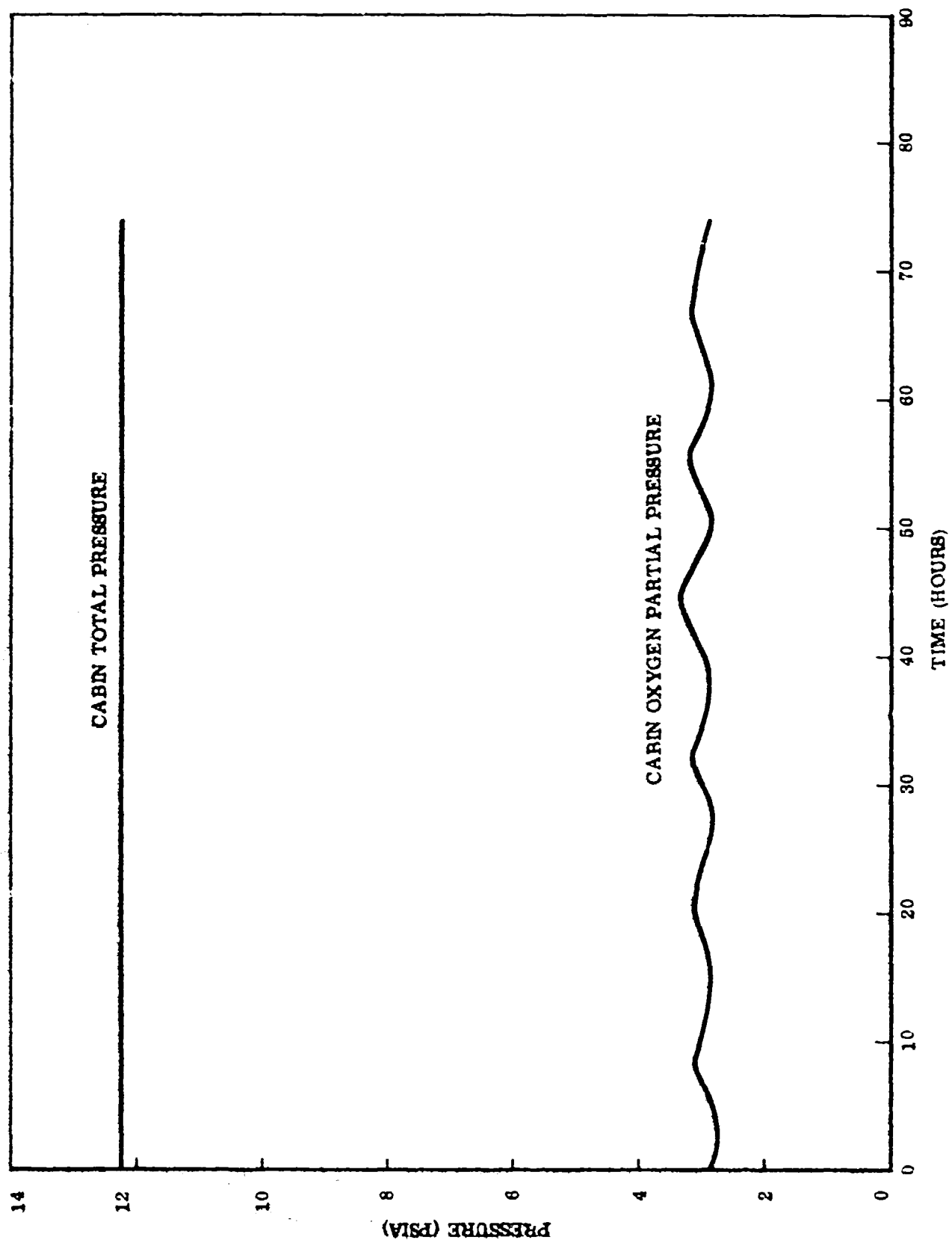


Fig. 2-32 Test 3 - Cabin Atmosphere Control

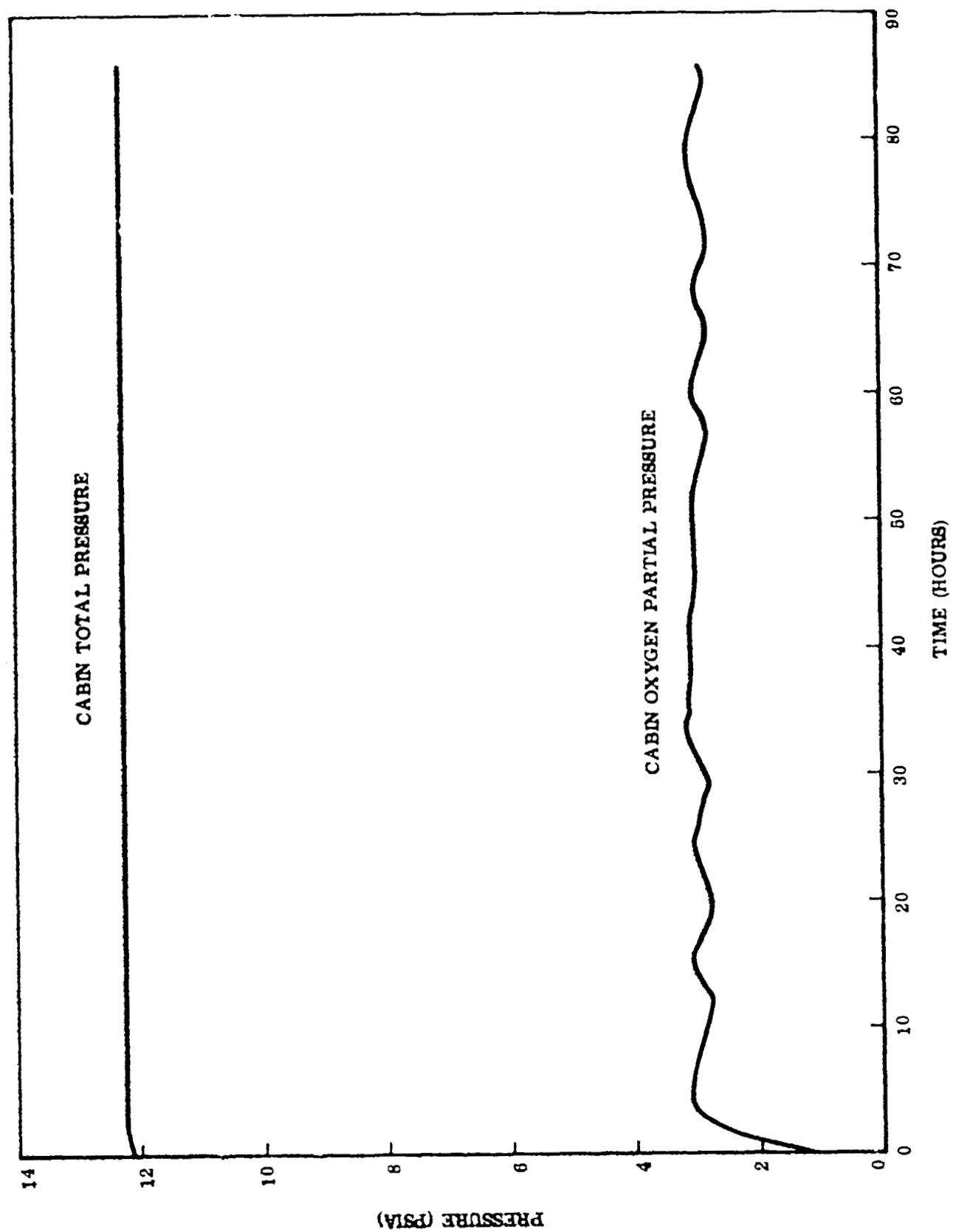


Fig. 2-33 Test 4 - Cabin Atmosphere Control

Table 2-3  
TEST 1 - TIME/EVENT LOG

Elapsed Time (hr)	Events/Actions	Criteria
8	Leak in the plumbing between the cell stack and the cabin simulator detected and corrected	High cabin pressure observed
18	N <sub>2</sub> H <sub>4</sub> flowmeter replaced	Range of flowmeter incorrect
54	Calibration of PO <sub>2</sub> sensors changed from 174 to 155 mm Hg	Calibration not previously corrected for 2 psig in sensor canister
78.8	Automatic safety shutdown	Water feed failed "on," and high-volume shutdown actuated
81	System returned to manual mode	
86	Test terminated	

At this point the reservoir piston had moved, because of the hydrazine addition, to a point where both the "on" and "off" water feed control microswitches were actuated. An error in the control logic permitted the "on" command to override the "off" command and resulted in continuous feed until the high reservoir volume signal shut down the system.

The system was restarted and operated for another 5 hr in a manual mode.

The following modifications to the system and test procedure were implemented as a result of this test:

- The PO<sub>2</sub> safety shutdown logic was modified to simply shut off the hydrazine feed solenoid valve, rather than shutting down the entire system. That is,

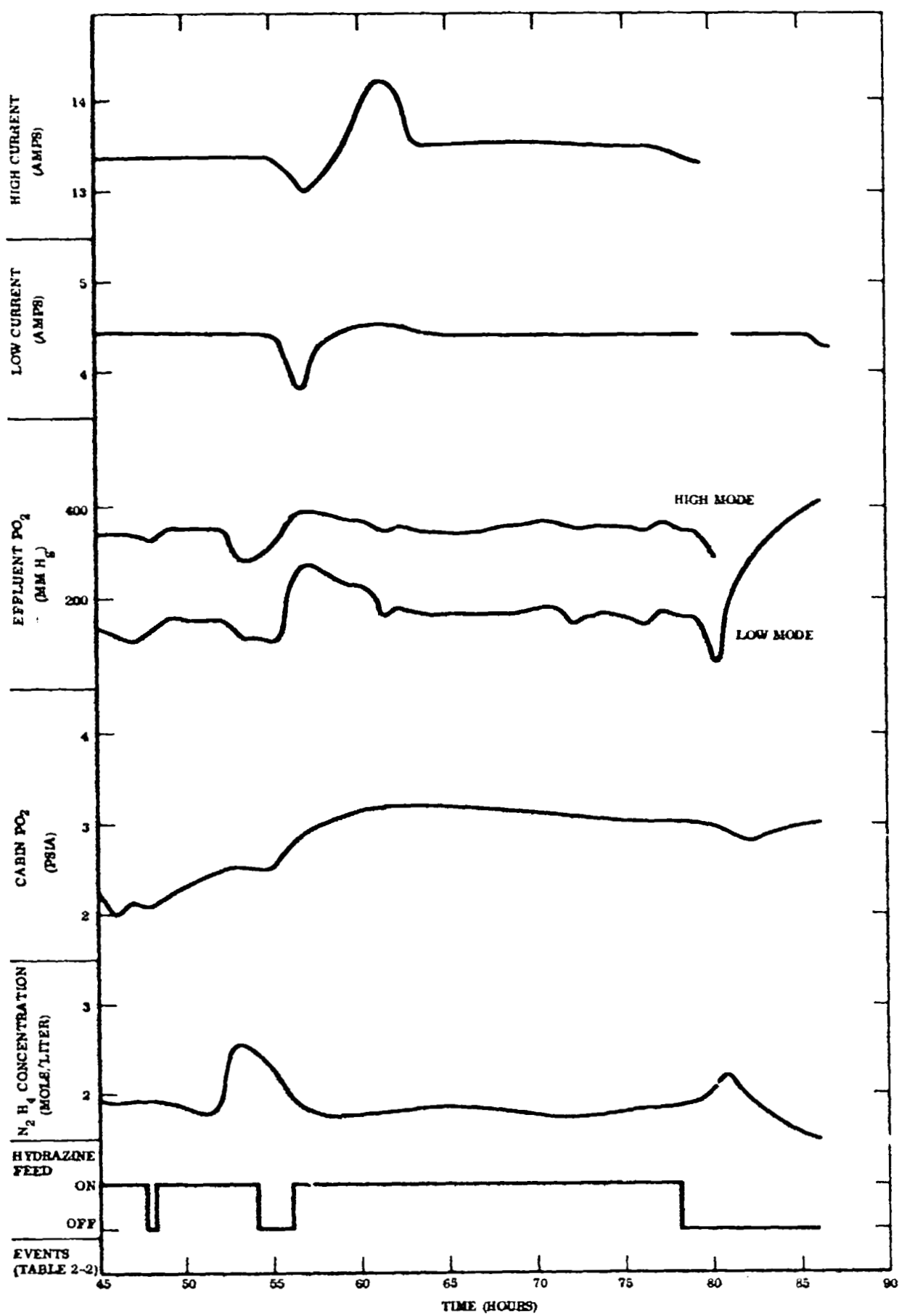


Fig. 2-34 Test 1 - Partial Plot of Performance

when the  $PO_2$  in the unit effluent decreased below 150 mm, the hydrazine feed would be stopped. This change resulted from observing a much faster hydrazine decay rate in the electrolyte than had been anticipated.

- A wet test meter was installed downstream of the metabolic/leak flowmeter to provide a more accurate measure of the gas removed from the cabin.
- The water feed control logic was corrected to assure positive shutoff in the event both "on" and "off" switches were actuated.
- The startup procedure was changed to include an initial hydrazine feed higher than the set point as a test expediency to shorten the test time required to achieve  $PO_2$  control.

Test 2 Results. An initial baseline run was conducted with the objective of determining the  $P_T$  and  $PO_2$  control characteristics at design conditions. The metabolic/leak and nitrogen makeup flow rates were set at the design point of a one-man metabolic load (1.84 lb/day) and a cabin leakage of one-twelfth of the spacecraft leakage rate of 13 lb/day at 10 psia. The  $P_{Total}$  control signal band was set for 12.23 to 12.25 psia, and the cabin  $PO_2$  control signal band was set for 2.8 to 2.9 psia. The high/low current modes were set approximately at 150/50 mA/cm<sup>2</sup>.

A performance plot and the Time/Event Log for the 70 hr of this test are shown in Fig. 2-35 and Table 2-4, respectively.

$P_{Total}$  is not shown on the figure; it remained within the control band for the entire test.

It can be seen in these performance data that the  $PO_2$  control and hydrazine concentration in the electrolyte were erratic; although not out of control.

During the first  $PO_2$  cycle, several upward adjustments of the hydrazine feed rate were made because the  $PO_2$  overshoot of the control band appeared excessive and the cycle time appeared to be too long. At 39 hr, the hydrazine tank ran empty, before the end of the feed cycle, and was recharged. The amount of hydrazine in the original

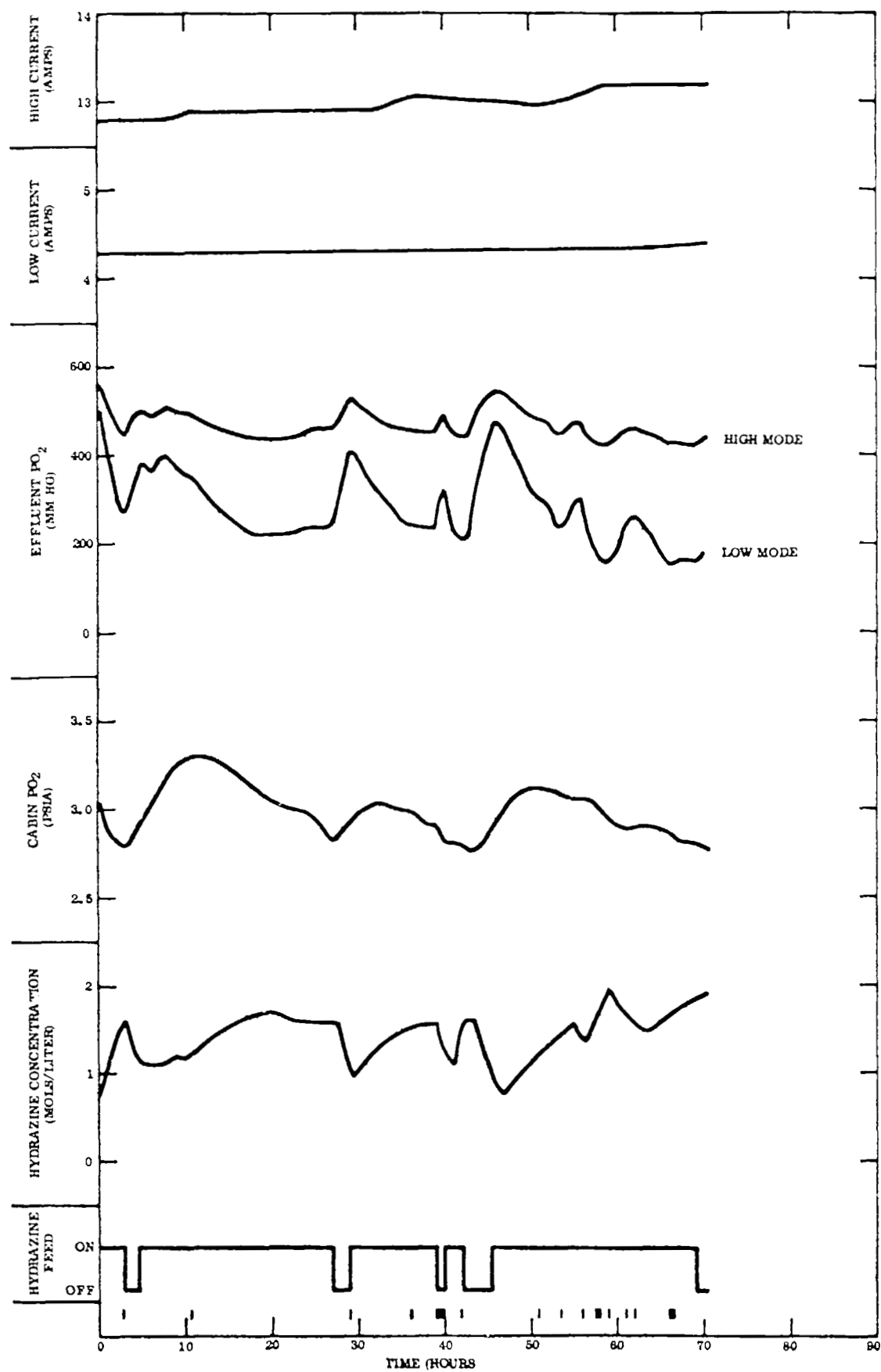


Fig. 2-35 Test 2 - Performance Data

Table 2-4  
TEST 2 - TIME/EVENT LOG

Elapsed Time (hr)	Events/Action	Criteria
2.7	Metabolic/leak and N <sub>2</sub> makeup set to correct settings for automatic control run; N <sub>2</sub> H <sub>4</sub> at 6.0 flowmeter reading	N <sub>2</sub> H <sub>4</sub> feed cycled off automatically
10.5	Adjusted N <sub>2</sub> H <sub>4</sub> to 7.0 flowmeter reading	Cabin PO <sub>2</sub> appeared to be overshooting, indicating N <sub>2</sub> H <sub>4</sub> feed rate too low
29	Adjusted N <sub>2</sub> H <sub>4</sub> to 7.5 flowmeter reading	N <sub>2</sub> H <sub>4</sub> feed cycled on and flow rate appeared low
36 to 36.2	N <sub>2</sub> makeup supply cylinder replaced	No-flow indication on flow control panel
39 to 40	Recharged N <sub>2</sub> H <sub>4</sub> tank; flow set high to restore hydrazine concentration in electrolyte	Tank ran empty
42	Adjusted N <sub>2</sub> H <sub>4</sub> to 5.8 flowmeter reading	Cabin PO <sub>2</sub> appeared to be decreasing too rapidly
51	Decreased coolant supply temperature to 35° F	Coolant to heat exchanger not cycling
53.5	Adjusted N <sub>2</sub> H <sub>4</sub> to 5.8 flowmeter reading	Cabin PO <sub>2</sub> leveled off
56	Adjusted N <sub>2</sub> H <sub>4</sub> to 7.7 flowmeter reading	
57.5 to 58.0	Repeated adjustment of N <sub>2</sub> H <sub>4</sub> downward to 6.5 from 7.6	
59	Adjusted N <sub>2</sub> H <sub>4</sub> to 6.0 flowmeter reading	
61	Adjusted N <sub>2</sub> H <sub>4</sub> to 6.5 flowmeter reading	
62	Adjusted N <sub>2</sub> H <sub>4</sub> to 7.0 flowmeter reading	
66 to 66.5	Adjusted N <sub>2</sub> H <sub>4</sub> to 6.9 flowmeter reading, then to 6.5	

charge had been predetermined based on the nitrogen load and was not expected to run out for another 12 hr. This incident shed doubt on the accuracy of the hydrazine flowmeter and led the test conductors to frequently adjust the hydrazine flow control valve setting during the remainder of the test in an effort to find the correct setting.

At the conclusion of the test it was still not clear why, during the first 40 hr of the test, a higher indicated flow rate was required than during the remainder of the test. Analysis of the hydrazine concentration in the solutions used to charge the tank revealed a discrepancy. In the first charge, bottles of hydrazine fresh from the manufacturer and labeled 95 percent were diluted with water by an amount which would make the desired solution 64 percent. Several bottles of this "95 percent" solution were analyzed and found to contain 64 percent solution. Because of the mislabeling, the actual solution had been diluted for charging the tank to approximately 40 percent. Further checking showed that the second charge of the tank was made with 64 percent solution.

Two problems with the hydrazine flow control were identified during the test, only one of which had an easy solution. First, the micrometer valve was too coarse and was replaced with a finer needle valve after the test was completed. Second, the valve body was made of 316 stainless steel, which was not completely compatible with hydrazine. A very slight decomposition of hydrazine inside the valve caused small gas bubbles to form in the flow stream. These small bubbles collected on the ball floats in the rotameter and gave rise to erroneous flow rate readings. The procedure for the next test was modified to include draining and recharging the hydrazine tank at the end of each feed cycle. By measuring the volume of the charge and the residual for each cycle, the consumption of hydrazine and the average feed rate for each cycle could then be determined.

Test 3 Results. The baseline operating conditions used in Test 2 were utilized in this test with the objective of obtaining a sufficient number of repeatable  $\text{PO}_2$  cycles to provide data for verifying the computer model revision. The only condition that was modified was the  $\text{PO}_2$  control signal band. It was changed from 2.8 - 2.9 psia to 2.85 - 3.00 psia as a result of recalibration of the sensors and the control electronics.



The performance data for Test 3 are shown in Fig. 2-36 and the Time/Events Log in Table 2-5.

During the first  $PO_2$  cycle, several operator adjustments of the hydrazine flow control valve were made in seeking the proper flow setting. After the first cycle, no further adjustments of the flow setting were made for the duration of the test.

The only problems that occurred during the test were two occasions when the hydrazine solenoid valve did not open on command. The first time this happened (at 6 hr elapsed time), the test conductor observed it immediately and was able to manually pulse the valve open. On the second occasion, however, the command and failure to open occurred at 41 hr and was not noticed until 42.7 hr. This resulted in the higher  $PO_2$  overshoot shown in Fig. 2-36 for the fourth cycle. With this exception, adequate  $PO_2$  control was demonstrated and reasonably repeatable cycles were obtained.

For the next test, the procedure for measuring hydrazine consumption was changed from a volume measurement to a weight measurement. This was done to increase the accuracy of the measurement and to reduce the handling required.

Test 4 Results. This test was run for a period of 86 hr with the objective of providing performance data at conditions different from the baseline. The performance data and Time/Event Log for this test are shown in Fig. 2-37 and Table 2-6, respectively.

During the first 29 hr of the test the baseline conditions were used, the only difference being that the low current was set at approximately  $75 \text{ mA/cm}^2$  rather than the baseline value of  $50 \text{ mA/cm}^2$ .

At 29 hr, the metabolic/leak and nitrogen makeup flow rates were changed corresponding to a change in a 12-man cabin from a 12-man to a 3-man metabolic load with no change in the cabin leakage.

After 45 minutes of operation under these conditions it was observed that cabin pressure,  $P_T$  was steadily increasing with the current in the low mode. This indicated

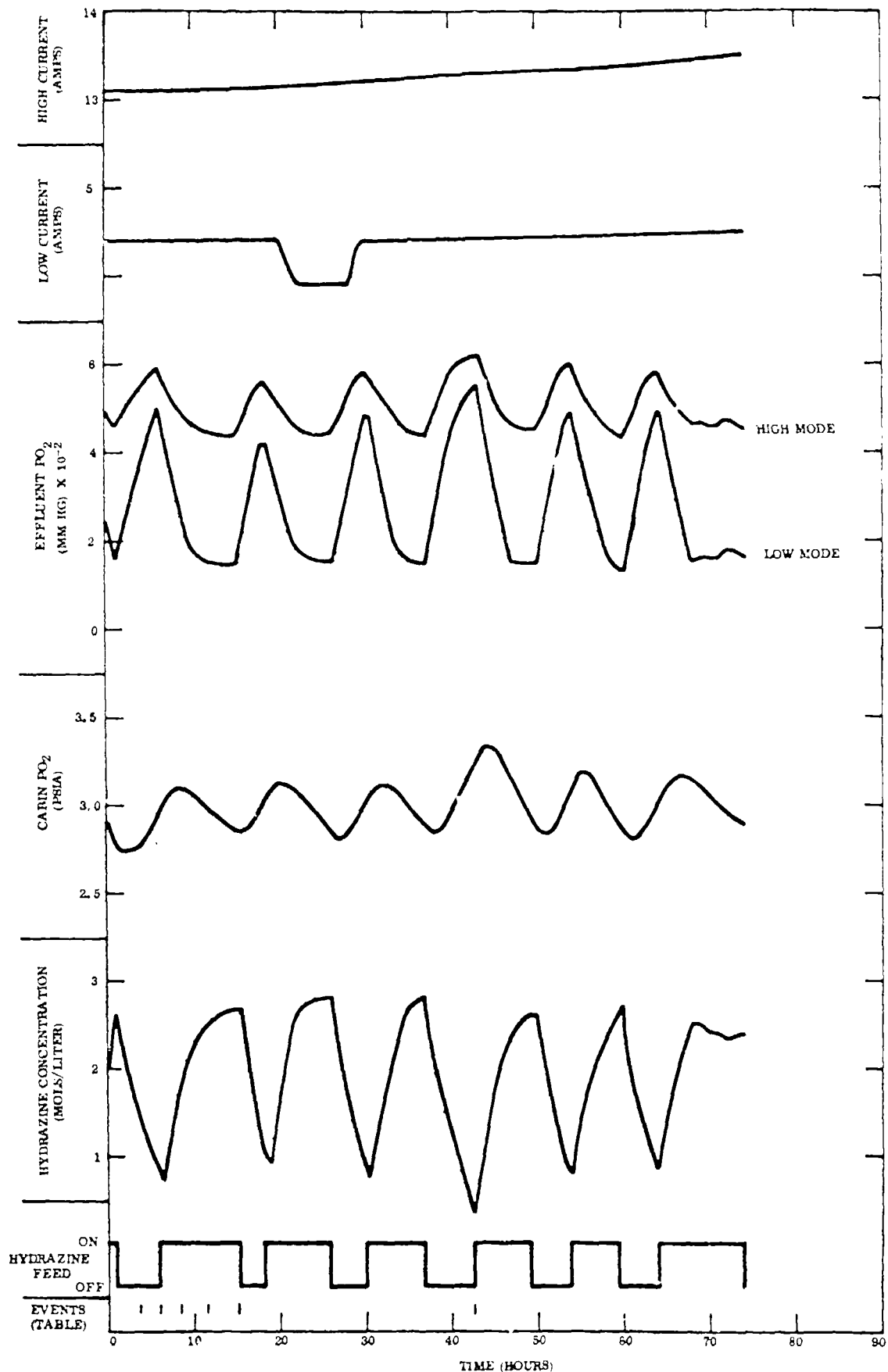


Fig. 2-36 Test 3 - Performance Data

Table 2-5  
TEST 3 - TIME/EVENT LOG

Elapsed Time (hr)	Events/Action	Criteria
1.1	Recharged N <sub>2</sub> H <sub>4</sub> tank; consumption on first cycle = 425 cc	Procedure
4.75	Adjusted metabolic/leak flowrate to 2,190 cc/min	Metabolic leak wet test meter reading low, at 2,110 cc/min
6	N <sub>2</sub> H <sub>4</sub> solenoid stuck closed; manually pulsed to open	N <sub>2</sub> H <sub>4</sub> feed signal on; no flow observed
8.5	Decreased N <sub>2</sub> H <sub>4</sub> flowmeter reading from 8.8 to 7.7	Noted electrolyte pressure increasing at approximately 1 in. H <sub>2</sub> O/hr
10	Increased N <sub>2</sub> H <sub>4</sub> flowmeter reading to 8.8	Noted electrolyte pressure dropping rapidly and effluent PO <sub>2</sub> climbing
11.5	Decreased N <sub>2</sub> H <sub>4</sub> flowmeter reading to 8.0	Effluent PO <sub>2</sub> safety signal
15.2	Decreased N <sub>2</sub> H <sub>4</sub> flowmeter reading to 7.9	Effluent PO <sub>2</sub> safety signal
15.5	Recharged N <sub>2</sub> H <sub>4</sub> tank; consumption on second cycle = 580 cc	Procedure
26	Recharged N <sub>2</sub> H <sub>4</sub> tank; consumption on third cycle = 510 cc	Procedure
37.3	Recharged N <sub>2</sub> H <sub>4</sub> tank; consumption on fourth cycle = 410 cc	Procedure
42.7	N <sub>2</sub> H <sub>4</sub> solenoid stuck closed; manually pulsed to open	Noted feed signal on; no flow indicated
50.5	Recharged N <sub>2</sub> H <sub>4</sub> tank; consumption on fifth cycle = 425 cc	Procedure
60.3	Recharged N <sub>2</sub> H <sub>4</sub> tank; consumption on sixth cycle = 370 cc	Procedure
End of test	Measured N <sub>2</sub> H <sub>4</sub> consumption on seventh cycle = 585 cc	Procedure
	Measured H <sub>2</sub> O consumption for duration of test = 2,330 cc	Procedure

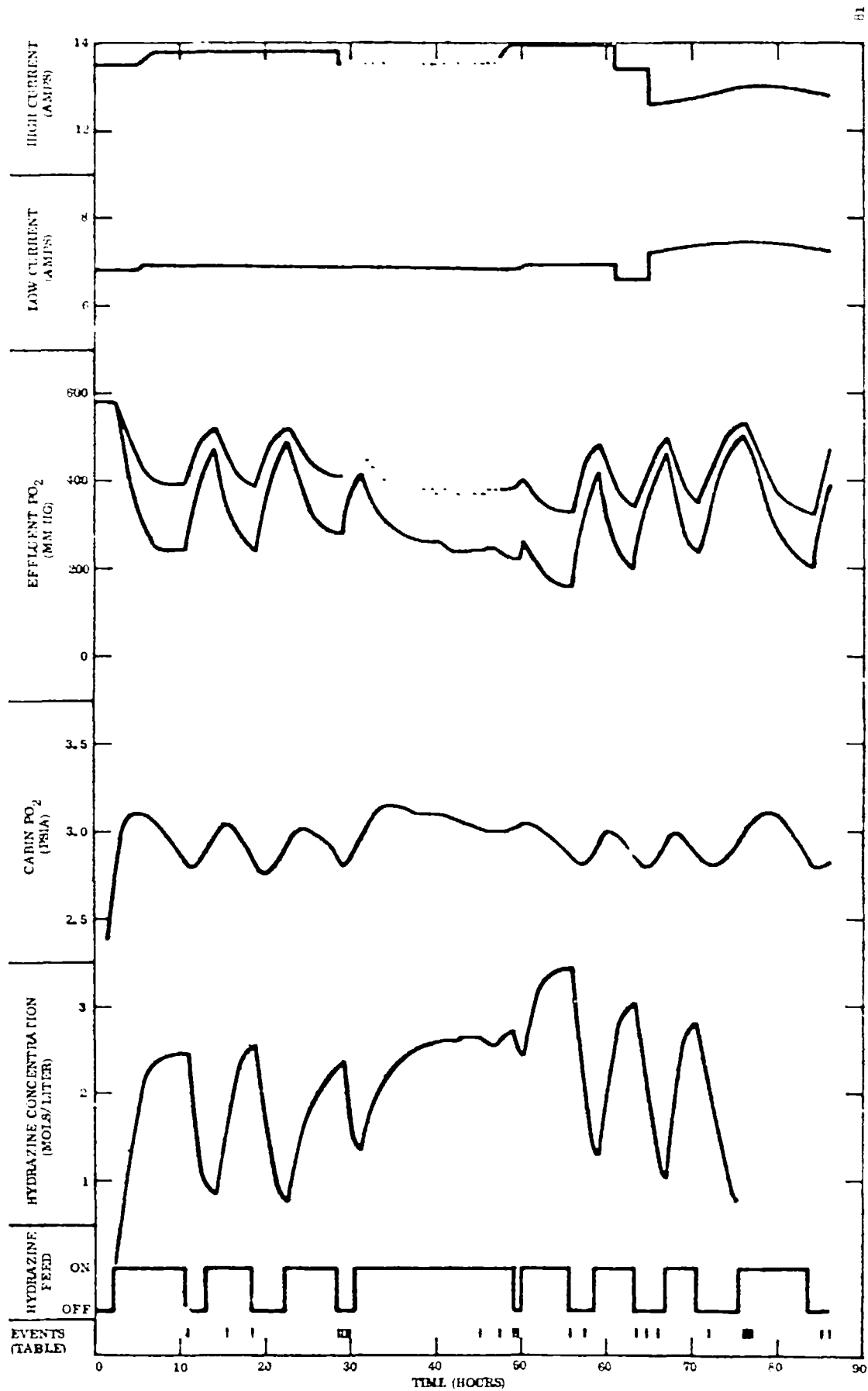


Fig. 2-37 Test 4 - Performance Data

Table 2-6  
TEST 4 - TIME/EVENT LOG

Elapsed Time (hr)	Events/Actions	
11	Recharged N <sub>2</sub> H <sub>4</sub> tank; consumption on first cycle = 575 gm	Procedure for measuring N <sub>2</sub> H <sub>4</sub> average feed rate
15.5	Adjusted N <sub>2</sub> H <sub>4</sub> feed rate to 3.93 valve setting	
18.5	Recharged N <sub>2</sub> H <sub>4</sub> tank; consumption on second cycle = 351 gm	Procedure
28.5	Adjusted high current down to 13.5 A	Current reading too high
29	Changed meta/leak and N <sub>2</sub> makeup from 12-man to 3-man metabolic rate	Test condition change
29.5	Recharged N <sub>2</sub> H <sub>4</sub> tank; consumption on third cycle = 441 gm	Procedure
29.75	Changed meta/leak and N <sub>2</sub> makeup to 4-man rate	Test condition change <sup>(a)</sup>
45.2	Changed to 5-man rate	Test condition change <sup>(a)</sup>
47.5	Changed to 6-man rate	Test condition change <sup>(a)</sup>
49	Recharged N <sub>2</sub> H <sub>4</sub> tank; consumption on fourth cycle = 1,299 gm	Procedure
49.5	Changed to 10-man rate	Test condition change
55.7	Changed to 12-man rate	Test condition change
57.5	Recharged N <sub>2</sub> H <sub>4</sub> tank; consumption on fifth cycle = 422 gm	Procedure
63.5	Adjusted high current to 13.5 A	Current too high
64.7	Changed currents: low mode 7.6 A; high mode 12.8 A	Test condition change
66	Recharged N <sub>2</sub> H <sub>4</sub> tank; consumption on sixth cycle = 320 gm	Procedure
72	Recharged N <sub>2</sub> H <sub>4</sub> tank; consumption on seventh cycle = 289 gm	Procedure
76 to 77	Shut off H <sub>2</sub> O and N <sub>2</sub> H <sub>4</sub> feed and inserted toggle valve in H <sub>2</sub> O feedline. Return both to auto feed mode	Indication by increasing system pressure that H <sub>2</sub> O solenoid leaking
85.2	Recharged N <sub>2</sub> H <sub>4</sub> tank; consumption on last cycle = 513 gm	Procedure
End of test	Measured H <sub>2</sub> O consumption for duration of test = 1,870 cc	Procedure

(a) System pressure rising

that the net flow of gas out of the cabin was less than the net gas flow in from the electrolysis module, and therefore, that  $P_T$  control could not be achieved under these conditions. Since an error of only 1 percent in the setting of the metabolic/leak flow rate could cause this problem, it was decided not to continue operation so close to this lower limit of  $P_T$  control.

At 29.75 hr, the metabolic/leak and nitrogen makeup were changed corresponding approximately to a four-man metabolic rate. This change restored  $P_T$  control and current-mode cycling was observed, although most of the time was being spent in low mode.

From 33 to 48 hr, a steady increase in electrolyte pressure was observed. During this period, electrolyte was manually removed from the system in 10-cc amounts approximately every half-hour to prevent the reservoir from reaching the high-volume shutdown position. Cause of the volume increase was first attributed to the increase in hydrazine concentration and decrease in water consumption resulting from the reduced metabolic load. Step-changes in the metabolic load were continued, going to a five-man rate at 45.2 hr, a six-man rate at 47.5 hr, a 10-man rate at 50 hr, and back to the baseline 12-man rate at 55.7 hr.

After the six-man rate was reached, the problem of increasing electrolyte volume appeared to have been solved.

At 64.7 hr, the high/low currents were changed as shown in Fig. 2-37.

A slow leak through the water feed solenoid valve was detected at 76 hr. A toggle valve was inserted and was operated manually for the remainder of the test. When the solenoid valve was disassembled at the end of the test, it was found that a piece of Teflon plumber's tape used to wrap the pipe threads on the fittings connecting to the valve had settled across the valve seat. This prevented complete closure of the valve and explains the increasing electrolyte volume that was observed during the run at the four- and five-man metabolic rates.

### 2.3.3.2 Data Analyses

Control Characteristics. The equation describing the variation of cabin total pressure with time can be expressed as (Ref. 7)

$$\frac{dP_T}{dt} = \frac{1}{\lambda T} \frac{dM_T}{dt} \quad (2.3)$$

where

$$\lambda T = \frac{V_C (MW)_T}{RT}$$

$$\frac{dM_T}{dt} = \dot{m}_{T_{in}} - \dot{m}_{T_{out}}$$

In these equations,  $\dot{m}_{T_{out}}$  represents the net mass outflow rate from the cabin, i.e., metabolic consumption plus cabin leakage. The total gas, oxygen plus nitrogen, supplied to the cabin by the  $O_2/N_2$  generator is represented by  $\dot{m}_{T_{in}}$  and is related to the number of cells,  $n$ , and the applied current,  $Y$ , by

$$\dot{m}_{T_{in}} = \gamma n Y \quad (2.4)$$

where  $\gamma$  is a constant.

Because the gas input rate is directly proportional to the current, two-position, high/low current control of  $P_T$  will have a cyclic characteristic and  $\frac{dP_T}{dt}$  will be linear.

An expression for the cabin oxygen partial pressure variation with time, similar to that for  $P_T$ , can be written as

$$\frac{dPO_2}{dt} = \frac{1}{\lambda O_2} \frac{dMO_2}{dt} \quad (2.5)$$

where

$$\lambda_{O_2} = \frac{V_C (MW)_{O_2}}{RT}$$

$$\frac{dM_{O_2}}{dt} = \dot{m}_{O_2 \text{ in}} - \dot{m}_{O_2 \text{ out}}$$

the net oxygen input rate,  $\dot{m}_{O_2 \text{ in}}$ , can be expressed in terms of  $O_2/N_2$  system parameters as

$$\dot{m}_{O_2 \text{ in}} = \gamma n Y - \alpha n C - \beta n C Y$$

where

$\alpha$  and  $\beta$  = constants

$C$  = hydrazine concentration in electrolyte

The hydrazine concentration in the electrolyte at a point in time ( $t$ ) can be expressed as

$$C = \frac{X}{m} + \left( C_o - \frac{X}{m} \right) \exp \frac{(-mnt)}{V} \quad (2.6)$$

where

$X$  = hydrazine feed rate

$m$  = reaction rate constant =  $f(Y)$

$C_o$  = hydrazine concentration at  $t = t_o$

$V$  = electrolyte volume



It is readily apparent that the control of cabin  $PO_2$  achieved by controlling the hydrazine feed rate,  $X$ , is not independent of the current which is used to control  $P_T$ .

To illustrate the characteristics of the control of  $P_T$  and  $PO_2$  and their interaction, a portion of the analog data from Test 3 was plotted on an expanded time scale and is shown in Fig. 2-38. First, it can be seen that the  $P_T$  control is cyclic, with a linear rate of change as predicted by the mathematical description.

Two characteristics of the  $PO_2$  control are evident in this figure. The effect of the exponential time function in the mathematical expression for rate of change of  $PO_2$  is apparent. Furthermore, it can be seen that the  $P_T$  cycles are superimposed on the  $PO_2$  curve.

The effluent  $PO_2$  curve on the far left of this figure, which is the oxygen partial pressure in the  $O_2/N_2$  mixture being generated in the cell stack, shows the effects of both the high/low current and the hydrazine feed. The rapid cycling of the effluent  $PO_2$  is the result of the high/low current cycling. The broad cycle of the envelope is the result of the hydrazine concentration in the electrolyte increasing when the hydrazine feed is on, and decaying when the feed is off.

Hydrazine Reaction Efficiency. Data from Test 3 were used to determine the hydrazine reaction efficiency, which is defined as the mol percent of the hydrazine reacting at the anode. The following is a summary of the methods of determination and the results:

- a. Nitrogen in the anodic effluent (determined by effluent  $PO_2$  sensor) was compared to the hydrazine feed (determined by measuring volume of  $N_2H_4$  consumed per cycle of the cabin  $PO_2$ ). This method yielded a hydrazine reaction efficiency of 63 percent.
- b. Nitrogen in the anodic effluent (above method) was compared to nitrogen in the cathodic effluent (determined by gas chromatographic analysis). This method yielded a hydrazine reaction efficiency of 59 percent.

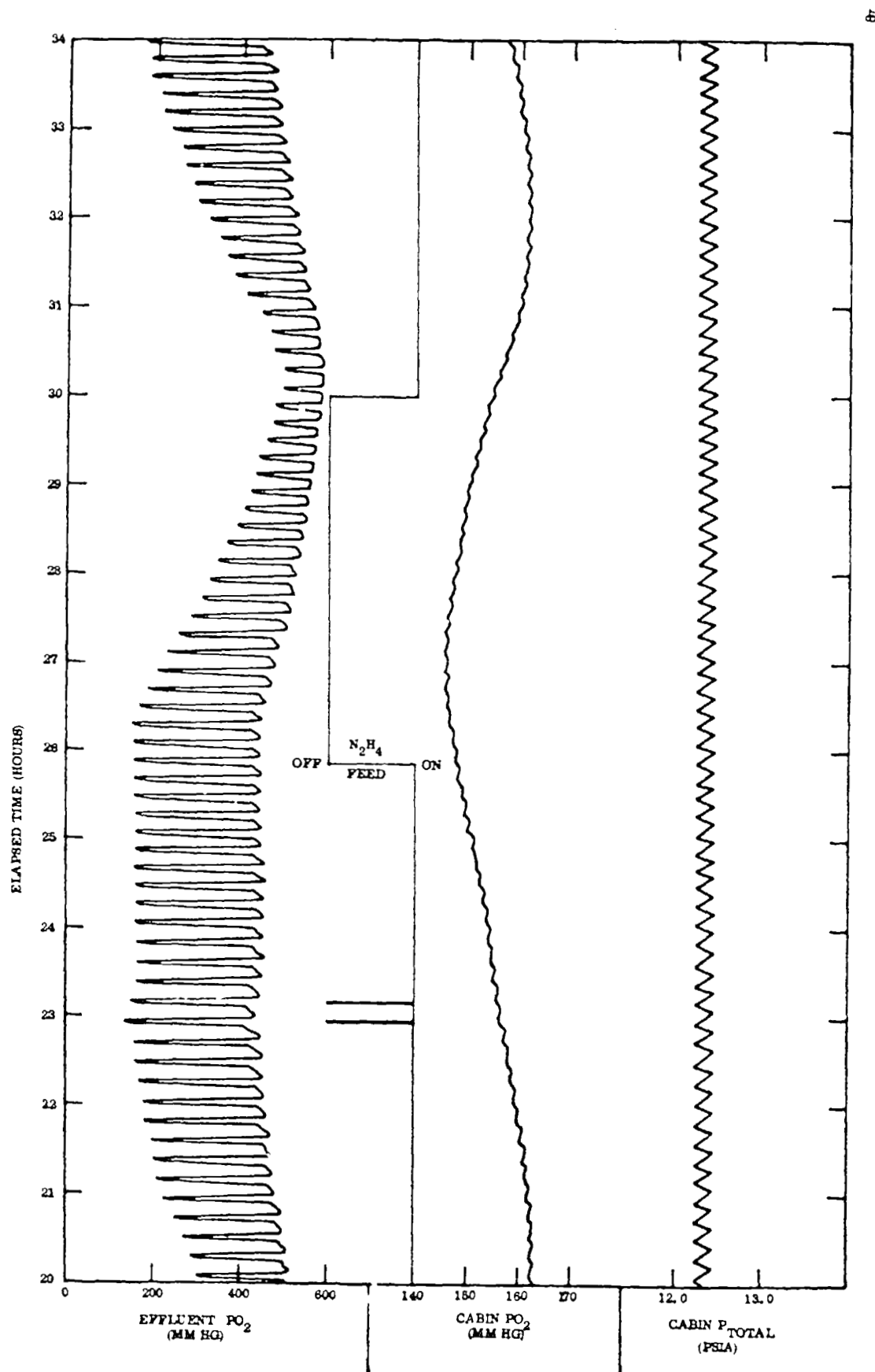


Fig. 2-38 Sample Data From Test 3

- c. Nitrogen in the cathodic effluent (determined by the increased cathodic flow-rate) was compared to the hydrazine feed determined in a. This method yielded a value of 62 percent.

Method b. involved point measurements whereas methods a. and c. utilized values averaged over a several-hour period. Details of the computations, data, and data references are given in Tables 2-7, 2-8, and 2-9. The results shown in these tables were averaged to obtain the efficiencies given above.

It should be noted that the inefficiency of the hydrazine reaction, which amounts to that hydrazine reacting undesirably at the cathode, is a function of the catalytic activity of the cathode. Single-cell tests have demonstrated that by sufficiently reducing the catalytic activity of the cathode, cathodic hydrazine reaction can be essentially eliminated. However, an increase in electrode polarization also occurs. Data from these tests are shown in Tables 2-10 and 2-11. In Table 2-10, the data were taken at a current density of  $50 \text{ mA/cm}^2$ , and in Table 2-11 at  $75 \text{ mA/cm}^2$ . Two separate cells were run, one containing the standard commercial (black platinum/Teflon) cathode and the second in which bare expanded nickel screen was used as the cathode with no platinum catalyst. From these limited data, it is apparent that the low catalytic activity cathode does not induce hydrazine decomposition. Note the lower cathodic flow rate for the low-activity cathode and the absence of  $\text{N}_2$  in the chromatographically analyzed cathodic gas samples.

Reaction Rate Constant. The hydrazine reaction rate constant is defined as

$$m = \frac{1}{C} \frac{dC}{dt}$$

and is truly a constant for a fixed cell geometry, temperature, and current. The effect of current at constant temperature on the value of  $m$  was determined using Test 3 data and is shown graphically in Fig. 2-39. Reducing the cathode catalytic activity to improve the hydrazine reaction efficiency would affect the value of the reaction rate constant.

Table 2-7

## HYDRAZINE CONVERSION EFFICIENCY - METHOD 1

Cycle	N <sub>2</sub> H <sub>4</sub> Feed (cc)	N <sub>2</sub> in Feed (mol)	Cycle Time	$\Delta t$ (hr)	N <sub>2</sub> in Anodic Effluent		Mode Time		Total Mode Current		N <sub>2</sub> in Anode Effluent (mol)	Efficiency (%)
					High (%)	Low (%)	High (hr)	Low (hr)	High (A)	Low (A)		
	(a)	(b)	(c)	(d)	(e)		(f)		(g)		(h)	(i)
1	580	11.6	6.17 to 18.1	11.93	27.4	63.4	8.04	3.89	224.5	75.64	6.56	56.5
2	510	10.2	18.1 to 30.1	12.0	32.0	59.7	7.57	4.43	225.8	71.53	7.06	69.1
3	410	8.2	30.1 to 42.75	12.65	24.7	52.3	8.04	4.61	226.2	75.72	6.08	74.3
4	425	8.5	42.75 to 53.4	10.65	24.5	63.2	6.26	4.39	227.9	74.30	5.34	62.8
5	370	7.4	53.4 to 63.95	10.55	23.5	55.6	6.64	3.91	226.2	74.28	4.93	66.7
6	585	11.7	63.95 to 73.94	9.99	28.1	65.7	6.97	3.02	228.4	74.94	5.73	49.1

(a) Volume of hydrazine consumed - Determined by measuring volume of initial charge and residual in hydrazine tank for each PO<sub>2</sub> cycle.

(b) Amount of nitrogen contained in Col. (a) - 64 percent N<sub>2</sub>H<sub>4</sub>  $\approx$  20 mole/liter  $\approx$  0.002 mole/cc; 1 mol N<sub>2</sub>H<sub>4</sub> contains one mole of cc; N<sub>2</sub> in feed = Col. a  $\times$  0.002.

(c) Cycle time - Times correspond to N<sub>2</sub>H<sub>4</sub> feed cycles (see Fig. 2-36).

(d)  $\Delta t$  - time interval between N<sub>2</sub>H<sub>4</sub> feed initiations.

(e) Percentage of N<sub>2</sub> in anodic effluent - Average N<sub>2</sub> concentration, determined by measuring the area under the PO<sub>2</sub> effluent curve (Fig. 2-36) for the high and low current modes for time periods (c), dividing by Col. (d) to obtain PO<sub>2</sub> avg, and computing the value of  $\%N_{2avg} = [(P_T - PO_{2avg})/P_T] \times 100$  where  $P_T \approx 12.23$ .

(f) High and low mode time - Determined by measuring total time on P<sub>T</sub> strip-chart recorder during which P<sub>T</sub> was increasing (equivalent to high current mode time) and the total time that P<sub>T</sub> was decreasing (equivalent to low current mode time).

(g) Total current, high and low mode - Total current in each mode, that is, average current during a cycle multiplied by the number of cells; determined by measuring the area under the current curve (Fig. 2-36) with a planimeter, dividing the area by (d) to obtain average current, and multiplying by 17, the number of cells.

(h) Number of mol of N<sub>2</sub> in anodic effluent during the cycle - determined by computing the value of  $\text{mol N}_2 = 0.0096 [(\text{Col. } e)_H (\text{Col. } f)_H + (\text{Col. } e)_L (\text{Col. } f)_L]$  where 0.0096 is the constant k converted from SCC/min to mol/hr.

(i) Hydrazine conversion efficiency - Determined by computing the value  $\eta(\text{efficiency}) = (\text{Col. h/Col. b}) \times 100$ .

Table 2-8

## HYDRAZINE CONVERSION EFFICIENCY - METHOD 2

Elapsed Time (hr)	I (A)	Percent N <sub>2</sub> (H <sub>2</sub> )	N <sub>2</sub> in H <sub>2</sub> (mol/hr)	Eff PO <sub>2</sub> (mm)	N <sub>2</sub> in O <sub>2</sub> (%)	N <sub>2</sub> in O <sub>2</sub> (mol/hr)	Efficiency (%)
	(a)	(b)	(c)	(d)	(e)	(f)	(g)
-0.2	12.9	6.8	0.359	510	22.1	0.465	56.5
1.27	4.3	13.3	0.310	160	75.8	0.533	63.3
1.38	13.1	7.3	0.399	480	26.7	0.572	58.9
7.0	13.1	4.3	0.210	570	13.0	0.279	57.0
16.13	13.3	7.9	0.449	500	23.6	0.512	53.3
30.25	13.2	3.1	0.148	600	8.4	0.181	55.1
37.33	13.4	8.7	0.514	450	31.2	0.683	57.1
59.73	13.4	7.4	0.415	430	34.3	0.751	64.5
60.0	4.5	14.4	0.372	130	80.2	0.590	61.3
65.7	13.4	5.4	0.282	530	19.0	0.416	59.6
65.95	4.4	12.9	0.302	290	55.7	0.400	57.1

(a) Current - Average cell current for the two cell stacks.

(b) Percentage of N<sub>2</sub> in cathodic effluent - gas chromatographic analysis.

(c) Nitrogen generation rate at cathode - determined by the following computation:

$$(H_2)_e = 2 \times 17 \times 0.0096 \times \text{Col. a}$$

$$\dot{m}_{TC} = (H_2)_e \times X_1 \times X_2,$$

where  $(H_2)_e$  = H<sub>2</sub> generated electrochemically (mol/hr)

$\dot{m}_{TC}$  = total gas generated at cathode (mol/hr)

$X_1$  = mol/hr N<sub>2</sub> from N<sub>2</sub>H<sub>4</sub>

$X_2$  = mol/hr H<sub>2</sub> from N<sub>2</sub>H<sub>4</sub> =  $2X_1$

$X_1$  = Col. b  $\times \dot{m}_{TC}$  = Col. b  $\times H_2 \times$  Col. c +  $3X_1$

$X_1$  = [Col. b  $\times (H_2)_e$ ]  $\div$  [1 - (3  $\times$  Col. b)]

(d) Effluent PO<sub>2</sub> - Instantaneous value of effluent PO<sub>2</sub> at the time the gas chromatograph sample was taken (see Fig. 2-28).

(e) Percentage of N<sub>2</sub> in anodic effluent - Determined by computing  $\%N_2 = (P_T - \text{Col. d}) / P_T$ .

(f) Nitrogen generation rate at anode - Determined by computing  $\dot{m}_{N_2} = 17(0.0096) \times \text{Col. a} \times \text{Col. e}$ .

(g) Hydrazine conversion efficiency - Determined by computing  $\eta = \text{Col. f} / (\text{Col. c} + \text{Col. f} \times 100)$

Table 2-9

## HYDRAZINE CONVERSION EFFICIENCY - METHOD 3

Cycle	Cathodic Gas Flow (cc)	Cathodic Gas Flow (mol)	$(H_2)_e$ (moles)	Efficiency (%)
	(a)	(b)	(c)	(d)
1	1,130	46.5	40.4	82.5
2	1,320	54.3	38.9	50.0
3	1,100	45.2	41.7	83.4
4	1,120	46.1	33.7	49.0
5	1,090	44.8	34.4	53.2
6	1,040	42.7	34.9	77.8

- (a) Cathodic gas flow (cc) - Total cathodic gas per cycle measured with wet test meter (See Appendix C)
- (b) Cathodic gas flow (mol) - Total cathodic gas per cycle converted from cubic centimeters to mols.
- (c) Electrochemically generated  $H_2$  - Determined from data in Table 2-8 by computing

$$(H_2)_e = 2 (0.0096) (I_{T_h} t_h + I_{T_l} t_l)$$

- (d) Hydrazine conversion efficiency - Determined by computing

$$\dot{m}_{T_C} = (H_2)_e + (X_1)_{N_2} + 2 (X_1)_{H_2}$$

$$X_1 = \dot{m}_{T_C} - (H_2)_e$$

$$\eta = \frac{N_2H_4 \text{ feed (mol)} - X_1}{N_2H_4 \text{ feed (mol)}} \times 100$$

Table 2-10

## HYDRAZINE CONVERSION - TEST 1

Standard Cell						Cell With Reduced Cathode Activity							
Time	Flow (cc/min)		Nitrogen Content (%)		N <sub>2</sub> H <sub>4</sub> Concen. (mol/ liter)	Cell Voltage (V)	Time	Flow (cc/min)		Nitrogen Content (%)		N <sub>2</sub> H <sub>4</sub> Concen. (mol/ liter)	Cell Voltage (V)
	O <sub>2</sub>	H <sub>2</sub>	O <sub>2</sub>	H <sub>2</sub>				O <sub>2</sub>	H <sub>2</sub>	O <sub>2</sub>	H <sub>2</sub>		
1:33					1.42	1.666	1:33					1.70	2.156
1:40	17.4						2:02		35.8				
1:57		46.2					2:07	17.9					
2:30					1.37							1.68	
2:38			43.2				2:33			53.7			
2:57	16.9	46.6					2:45			50.3			
2:53			42.4				3:02						
3:11				9.6			3:15	17.8	35.5				
3:29				9.6		1.668	3:19				0		
3:30							3:30				0		2.192
4:12			27.6				4:03			46.4			
4:18				8.8			4:28				0		
4:20	17.2	46.3					4:30	17.9	35.3				
Decomposition Rate (No Current) O <sub>2</sub> Side - 10.9 cc/min H <sub>2</sub> Side - 16.7 cc/min						Decomposition Rate (No Current) O <sub>2</sub> Side - 12.6 cc/min H <sub>2</sub> Side - None							

Table 2-11

## HYDRAZINE CONVERSION - TEST 2

Standard Cell							Cell With Reduced Cathode Activity						
Time	Flow (cc/min)		Nitrogen Content (%)		N <sub>2</sub> H <sub>4</sub> Concn. {mol/ liter}	Cell Voltage (V)	Time	Flow (cc/min)		Nitrogen Content (%)		N <sub>2</sub> H <sub>4</sub> Concn. {mol/ liter}	Cell Voltage (V)
	O <sub>2</sub>	H <sub>2</sub>	O <sub>2</sub>	H <sub>2</sub>				O <sub>2</sub>	H <sub>2</sub>	O <sub>2</sub>	H <sub>2</sub>		
11:14						1.807	11:18						2.324
11:15		70.0					11:31	26.3	51.6			1.99	
11:26	26.3						11:45						
11:36			37.6				12:11			37.9			
11:45					1.97		12:23			38.9			
11:48			37.9				12:47				0		
12:00			38.2				1:09					1.97	
1:07					1.80		1:50	26.5	53.1	37.3			
1:40			36.5				2:16				0		
1:45	26.2	67.6					2:35			37.3			
1:58				7.8									
2:04				7.6									
2:25			36.2										



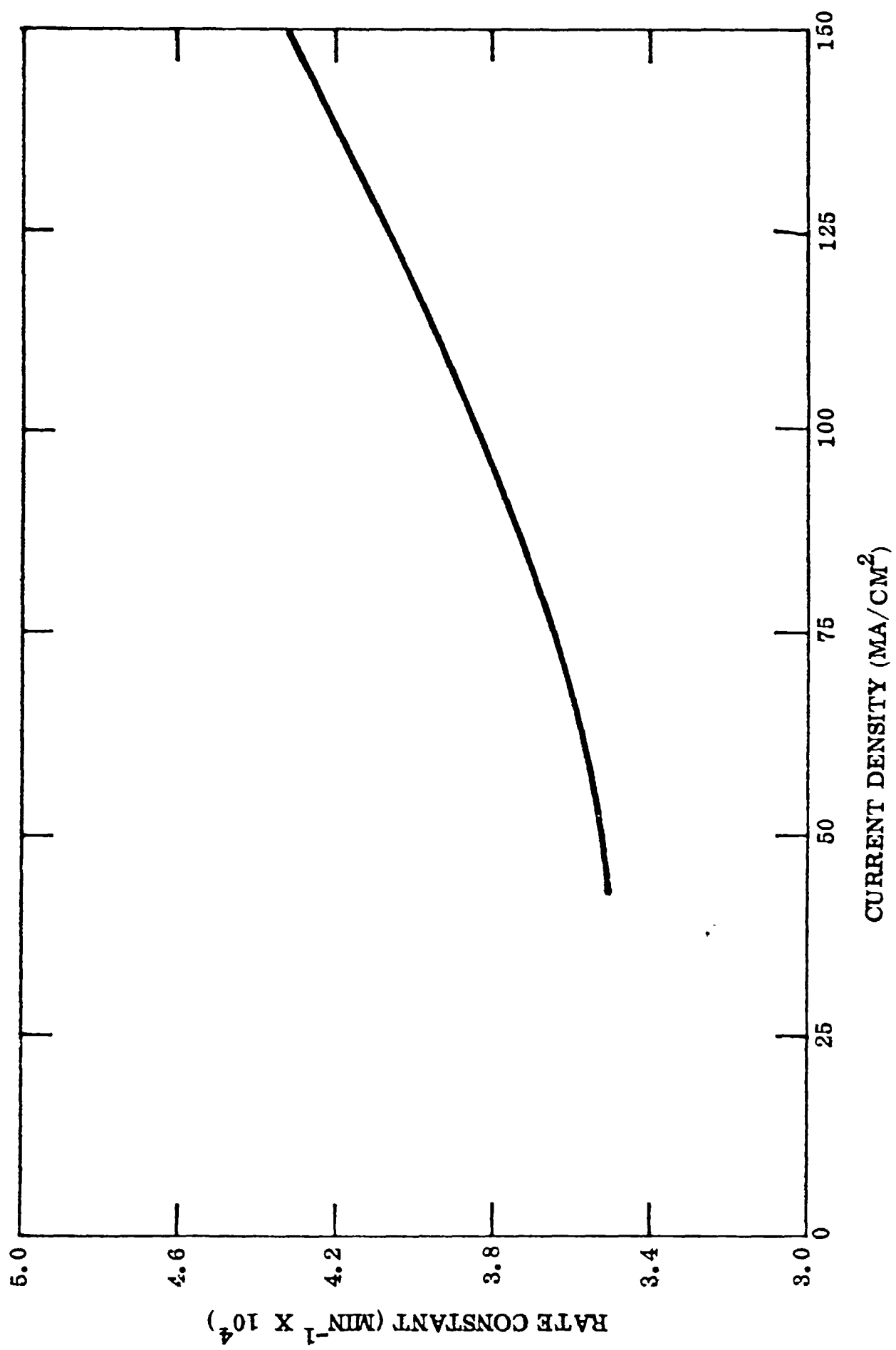


Fig. 2-39 Reaction Rate Constant vs. Current Density

Computer Model Constants. The experimentally determined constants required for the computer model updating, which is described in Section 2.4, were obtained by averaging several data points from Test 3. The analysis is described below.

The volumetric anodic gas generation rate, oxygen plus nitrogen, can be expressed as

$$\dot{U}_T = k n Y = k n \frac{I}{A}$$

where

$\dot{U}_T$  = total gas production, cc/min

$k$  = 0.35 (constant)

$n$  = 17 (number of cells)

$Y$  = current density, mA/cm<sup>2</sup>

$I$  = current amps

$A$  = 90 (electrode area, cm<sup>2</sup>)

Therefore:

$$\dot{U}_T = \frac{(0.35)(17)(10^3)}{90} I = 66.1 I$$

That portion of  $\dot{U}_T$  that is nitrogen can be expressed in terms of the polarographically measured effluent  $PO_2$  and the total pressure of the effluent gas as

$$\dot{U}_{N_2} = \left( \frac{P_T - PO_2}{P_T} \right)_{\text{effluent}} \dot{U}_T \quad (2.7)$$

At the point in the system where effluent  $PO_2$  is measured, the effluent gas pressure is the same as the cabin total pressure.

The empirical formula for  $\dot{U}_{N_2}$  is of the form

$$\dot{U}_{N_2} = a n C + b n C Y \quad (2.8)$$

or

$$\dot{U}_{N_2} = a n C + b n C \frac{I}{A} \quad (2.9)$$

Inserting known values, this becomes

$$\dot{U}_{N_2} = 17 a C + 189 b C I \quad (2.10)$$

Combining the two equations for  $\dot{U}_{N_2}$  results in the expression

$$66.1 I \frac{P_T - PO_2}{P_T} = 17 a C + 189 b C I \quad (2.11)$$

Because the hydrazine concentration does not change rapidly with time while the current is cycling every few minutes, as was shown in Fig. 2-38, it is possible to make a hydrazine concentration determination and record a set of both high and low currents and high and low  $PO_2$  effluents essentially at that concentration. The values of the constants  $a$  and  $b$  can then be determined by solving a simple set of two simultaneous equations in two unknowns.

Several data points were taken during Test 3 in the manner described above. The experimental data and the resulting values of the constants are shown in Table 2-12.

The averages of these values are:

$$a = 4.77$$

$$b = 0.0134$$

Table 2-12  
EXPERIMENTAL CONSTANTS DETERMINATION

Elapsed Time (hr)	$C(a)$	Current, I (A)		Partial Pressure, $PO_2$		Constants	
		High	Low	High	Low	a	b
0	2.00	12.9	4.30	490	240	4.67	0.0129
6.28	0.78	13.1	4.25	590	500	4.23	0.0153
15.9	2.15	13.3	4.45	470	210	4.88	0.0128
18.2	0.99	13.0	4.5	570	440	5.36	0.0089
37.5	2.17	13.4	4.55	440	160	5.26	0.0177
40.9	0.76	13.25	4.35	590	490	4.87	0.0118
60.0	2.75	13.2	4.5	420	130	4.30	0.0163
64.14	0.83	13.35	4.35	590	490	4.59	0.0114

(a) Concentration of hydrazine.

#### 2.3.3.3 System Analysis

$O_2/N_2$  Generator System. In general, the performance of this system was satisfactory, as evidenced by the performance data presented previously in this section. However, some problem areas were identified as a result of system testing.

The current regulators that were used to condition 28 Vdc power and provide constant current to the electrolysis cell stack showed a tendency to drift, especially in the high current mode. These are prototype devices that were designed under Contract

NAS1-9728. Their performance could be improved by incorporating temperature compensation and additional filtering.

The performance of the prototype bubble separator used in the system was not satisfactory. By the end of Test 2, the pressure drop across the hydrophilic membrane had increased to the point where the electrolyte flow rate was becoming marginal. The unit was then rebuilt with new membranes before proceeding with the system testing. Although clogging of the membrane with particulate matter may be a partial explanation of the problem, it was also observed, with fresh membranes installed, that the initial pressure drop across the membrane was sufficient to cause nucleation of gas bubbles on the downstream side. While this may only be a materials problem, concurrent effort in a separate program is being directed to a new bubble separator design concept.

Controls. The cabin  $PO_2$  control technique that was tested during this program employs a fixed hydrazine input rate during the portion of the control cycle where nitrogen addition to the cabin is called for. When the cabin  $PO_2$  reaches a minimum set point, the hydrazine feed is shut off.

The hydrazine feed rate is set so that the  $PO_2$  in the effluent  $N_2/O_2$  mixture from the electrolysis module will not be driven below a set minimum. Because of this limitation on hydrazine input rate, the cabin  $PO_2$  overshoots the control band during the long period of time (usually several hours) required to increase the hydrazine concentration in the electrolyte sufficiently to effect  $PO_2$  turnaround. Furthermore, it was found to be difficult to accurately set and maintain the very low flow rate (less than 1 cc/min) required. Because of the modular design concept of the full-scale system, this limitation would remain in the larger system.

Under steady-state conditions, the control of cabin  $PO_2$  achieved in the model system with this technique was acceptable: A typical cycle of cabin  $PO_2$  at a set control band of 2.85 to 2.95 psia in Test 3 took approximately 8 hr with an overshoot to approximately 3.1 psia. However, this control is sluggish and may be a limiting factor in the ability of the system to respond to sudden changes in demand.

A modification in the control technique in which the hydrazine feed rate is increased to allow a fast ramp of the nitrogen output with two additional set points that cycle the hydrazine feed off at minimum effluent  $PO_2$  levels corresponding to the high and low current modes would affect a faster turnaround of cabin  $PO_2$  and substantially reduce the overshoot. The system also would then be less sensitive to the hydrazine feed rate, and extreme accuracy of the flow rate setting would not be necessary.

The results of attempting to operate the system with the total gas demand very close to the minimum total gas generation rate in Test 4 were described previously. In this case, it was attempted to operate with a three-man metabolic load and the baseline cabin leakage. Under these conditions, the cell stack was generating gas, in the low current mode, at 459 cc/min and the metabolic/leak and nitrogen makeup were supposed to be set at 801 and 333 cc/min, respectively. This meant that the net gas out was supposed to be 468 cc/min, or only 9 cc/min more than the generated gas rate.

From these numbers, it can be seen that only a 1-percent error in metabolic/leak flow setting could cause loss of control; and further, that a combined error in metabolic/leak, nitrogen makeup, and cell current of 2 percent would have the same effect. In future tests it would be advisable to incorporate micrometer flow controls and very accurately calibrated flow indicators for high- or low-limit operation.

## 2.4 COMPUTER ANALYSES

Experimental data from the one-man model system testing were used to revise and update the computer routine model of the system developed previously under NASA Contract NAS1-7706. In order to verify the computer routine, the program was run using average simulated test conditions, and the results were then compared with the actual test performance. Parametric modeling was performed of the hydrazine feed rate, the gas consumption, metabolic demand, and high/low current modes. These efforts are described in this section.

It is noted that the original computer routine and its revision and updating have been documented separately in a Utilization Manual. Therefore the complete program listing and case inputs are not included in this report.

### 2.4.1 Model Revision

The empirical functions which appear in the computer routine model, their original values, their revised values, and the equation in which they appear are shown in Table 2-13.

The values of  $\alpha$  and  $\beta$  were obtained by applying a density conversion to the experimentally determined values of  $a$  and  $b$  given previously in Section 2.3.3.

The value of  $\gamma$  was obtained by applying a density conversion to the value of  $K$ . The value of  $K$  was revised to account for the water vapor in the gas and approximately a 1-percent shunt current.

The values of  $m$  were taken from the experimentally determined relationship of  $m$  to current density shown in Fig. 2-39.

Hydrazine Feed Rate. The effect of hydrazine feed rate on  $PO_2$  control is shown in Figs. 2-40 and 2-41. The hydrazine feed rate in the first case is  $8.35 \times 10^{-4}$  mols/min/cell and  $1.16 \times 10^{-3}$  mols/min/cell in the second case. These figures illustrate

Table 2-13

## COMPUTER MODEL REVISIONS

Function	Relation	Initial Value	Revised Value
$\alpha$	$\alpha = \rho a$	$4.25 \times 10^{-3}$	$5.33 \times 10^{-3}$
$\beta$	$\beta = \rho b$	$2.43 \times 10^{-5}$	$1.55 \times 10^{-5}$
$\gamma$	$\gamma = \rho k$	$4.35 \times 10^{-4}$	$4.13 \times 10^{-4}$
$m_1$	$m_1 = f(Y_1)$	$3.28 \times 10^{-4} @ 150$	$4.32 \times 10^{-4} @ 150$
$m_2$	$m_2 = f(Y_2)$	$2.52 \times 10^{-4} @ 50$	$3.52 \times 10^{-4} @ 50$

Equations:

$$C_{\max (1)} = \frac{\gamma Y_1}{\alpha + \beta Y_1}$$

$$C_{\max (2)} = \frac{\gamma Y_2}{\alpha + \beta Y_2}$$

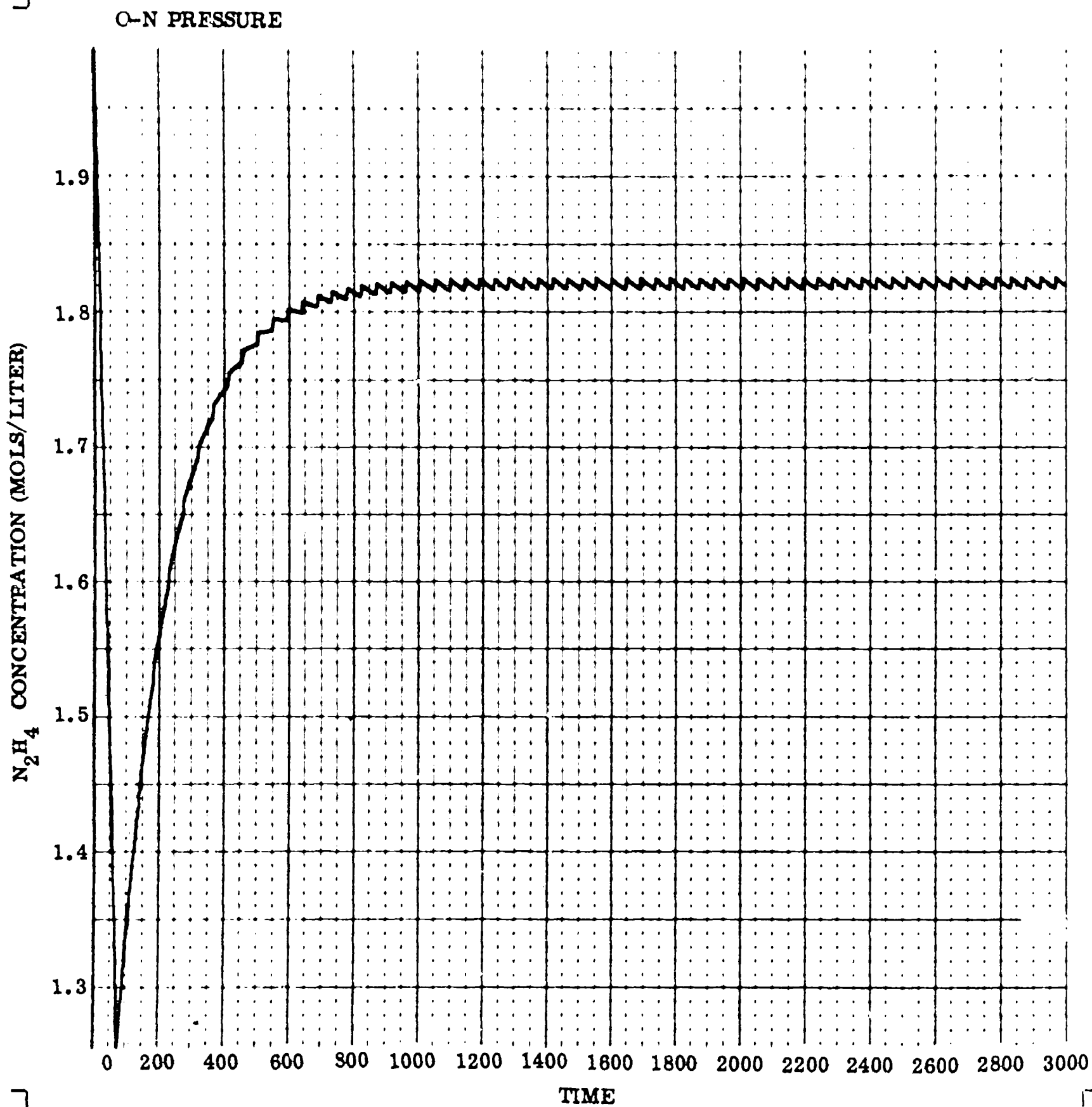
$$PO_2 = \frac{\lambda O_2 PO_{2i} + \left[ (\gamma n \cdot Y) - (\alpha n \cdot C) - (\beta n \cdot C \cdot Y) - \left( \dot{m}_{O_2 \text{ out}} \right) \right] \Delta t}{\lambda O_2}$$

$$P_T = \frac{P_{T_i} \lambda T + \left[ (\gamma n \cdot Y) - \left( \dot{m}_{T \text{ out}} \right) \right] \Delta t}{\lambda t}$$

$$C = \frac{X}{m} + \left( C_o - \frac{X}{m} \right) \exp (-mn t/v)$$

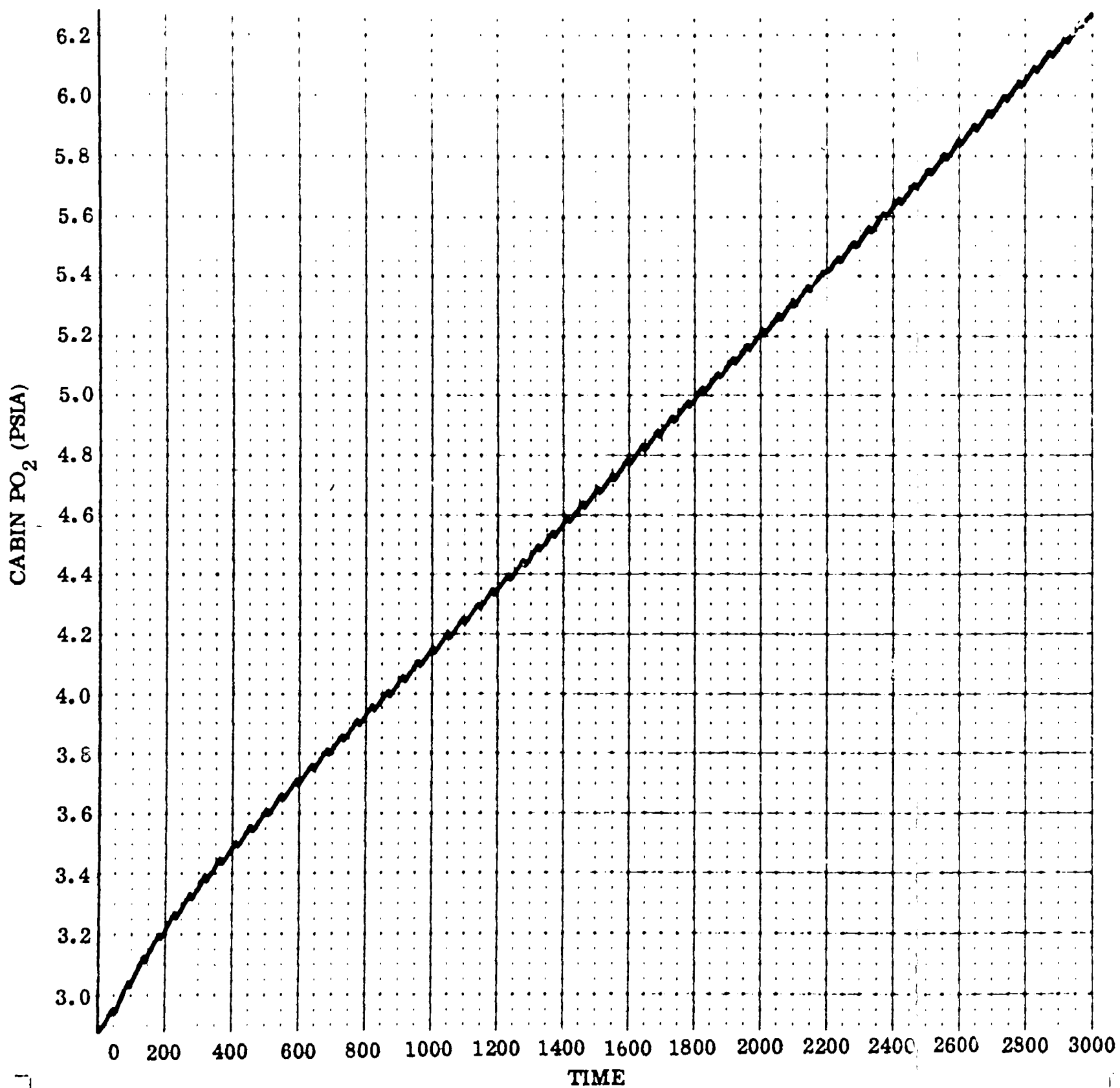
the critical effect of hydrazine flow rate on the  $PO_2$  control. In Fig. 2-40, the hydrazine concentration reaches an equilibrium value of 1.82 M, and insufficient nitrogen is being generated to effect turnaround of  $PO_2$ . At the higher hydrazine feed rate, in Fig. 2-41, the hydrazine concentration reaches 2.53 M, where sufficient nitrogen is being produced to cause the  $PO_2$  to cycle. The rate constants had not been revised at the time of these computer runs and therefore the cycle time shown in Fig. 2-41 is not representative.



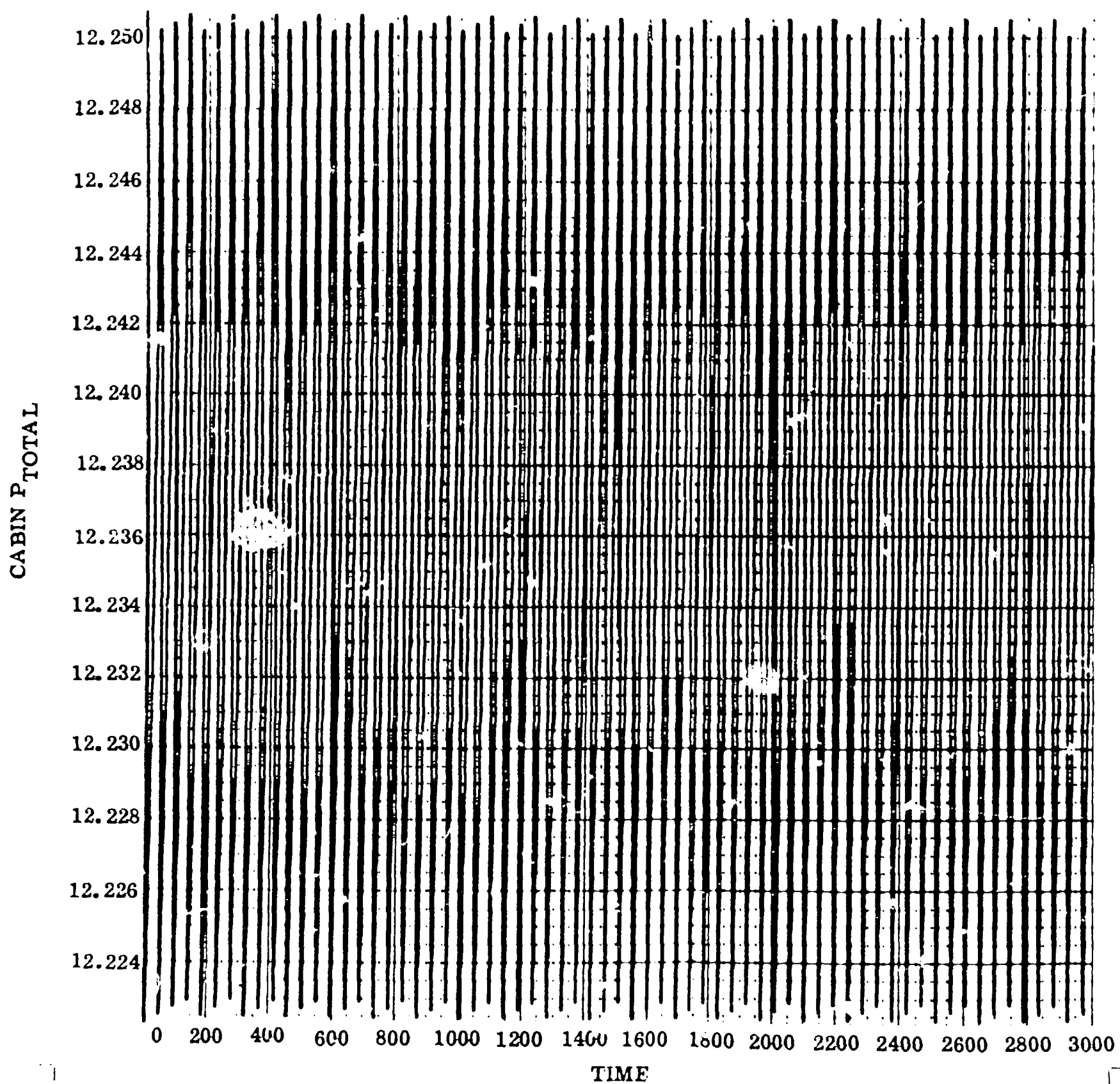


FOLDOUT FRAME

O-N PRESSURE



O-N PRESSURE



# O-N PRESSURE

CELLS: 20, 40, 60  
TIME: 1000

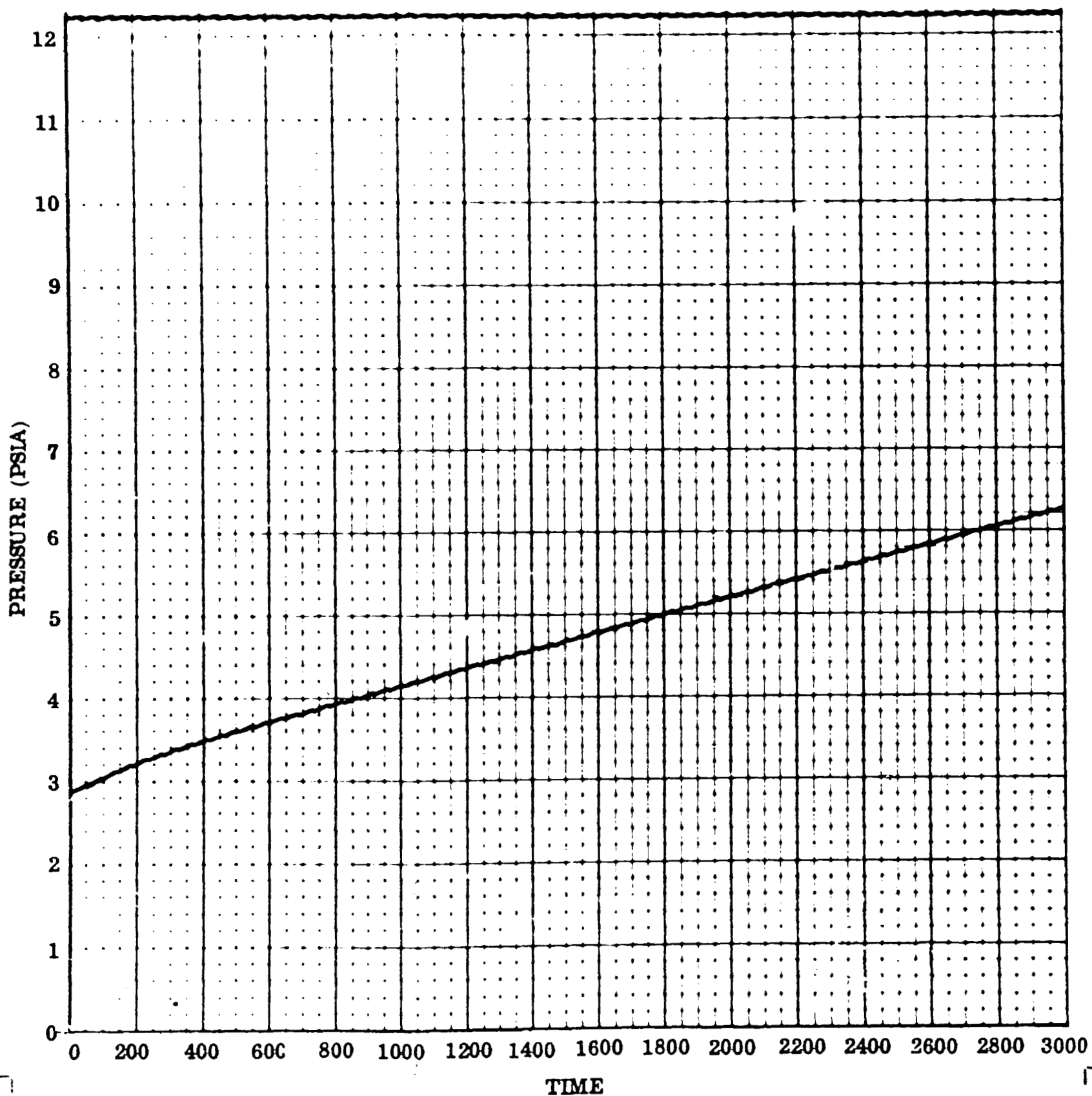
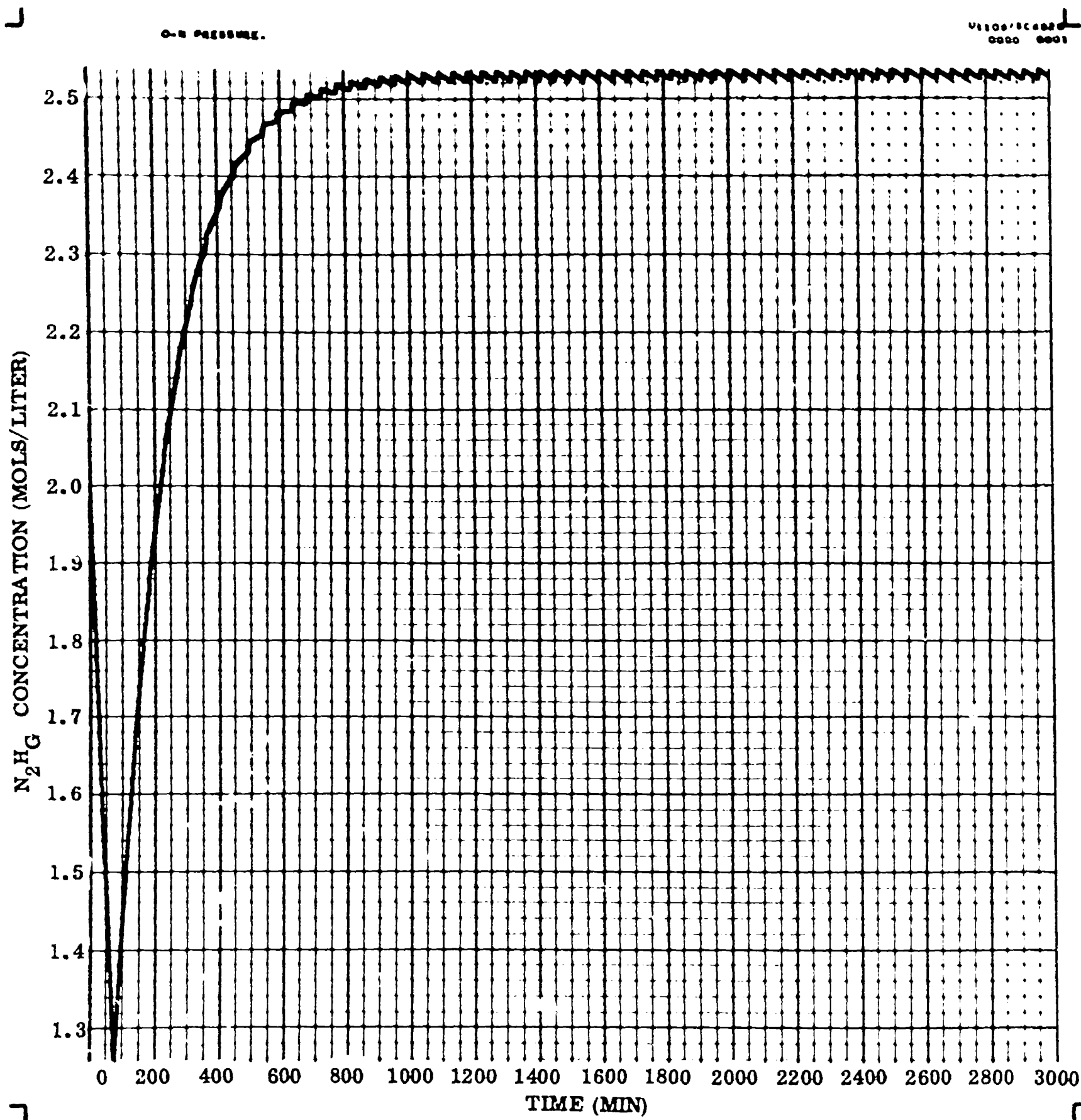


Fig. 2-40 Initial Run of Revised Computer Routine

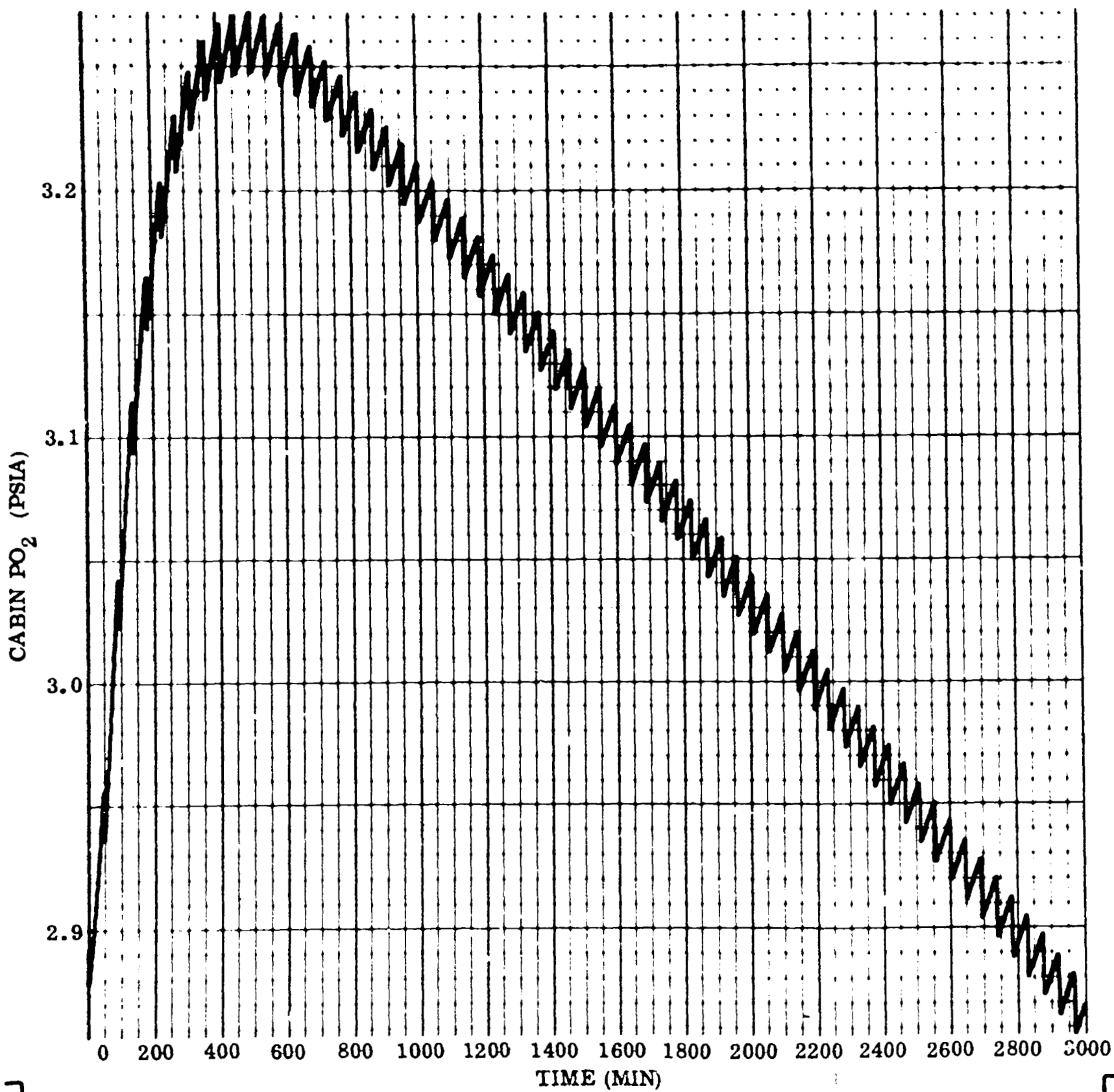


FOLDOUT FRAME 1

L

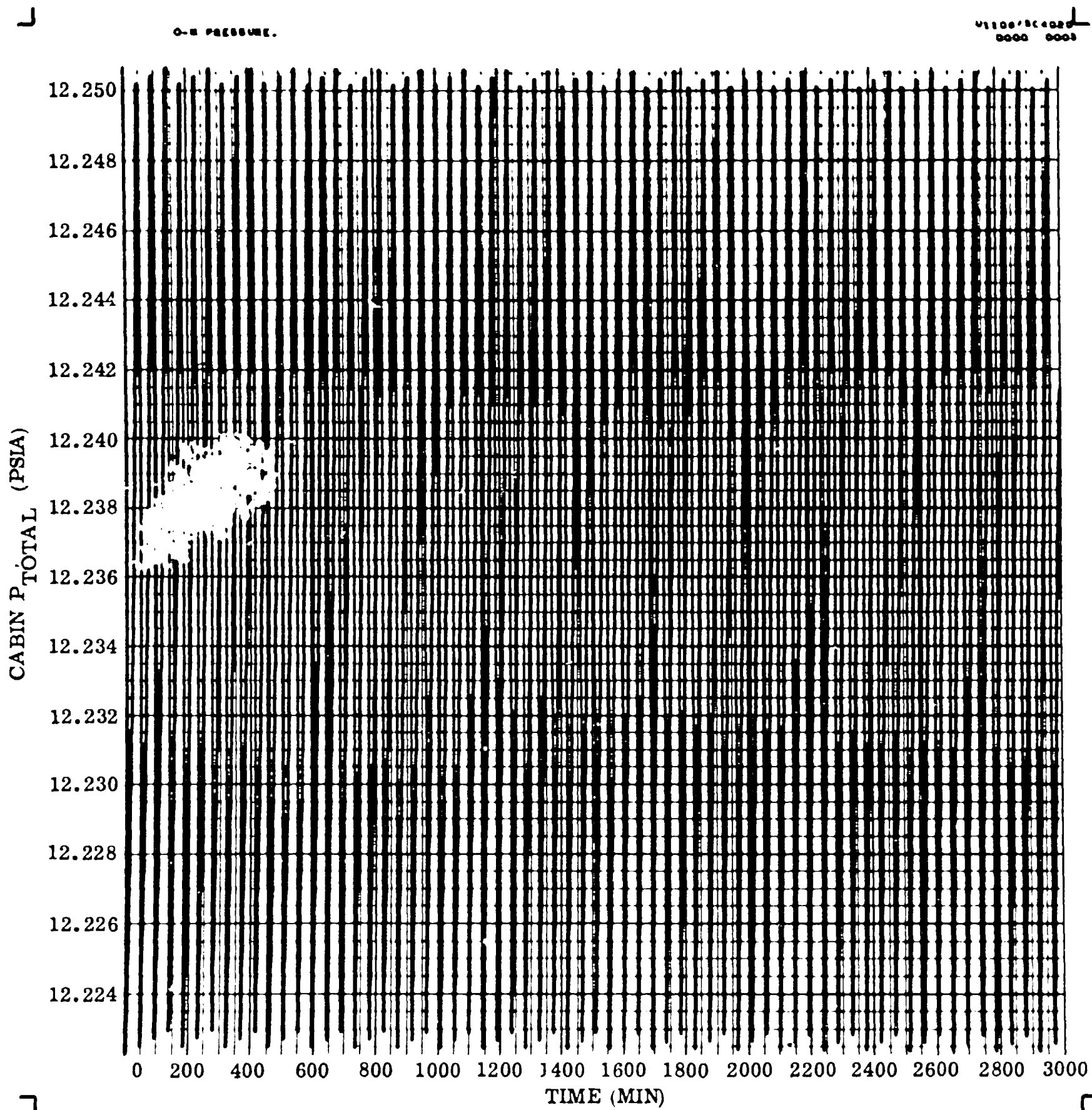
O-N PRESSURE.

VIDEO RECORDED  
0000 0000



J

r



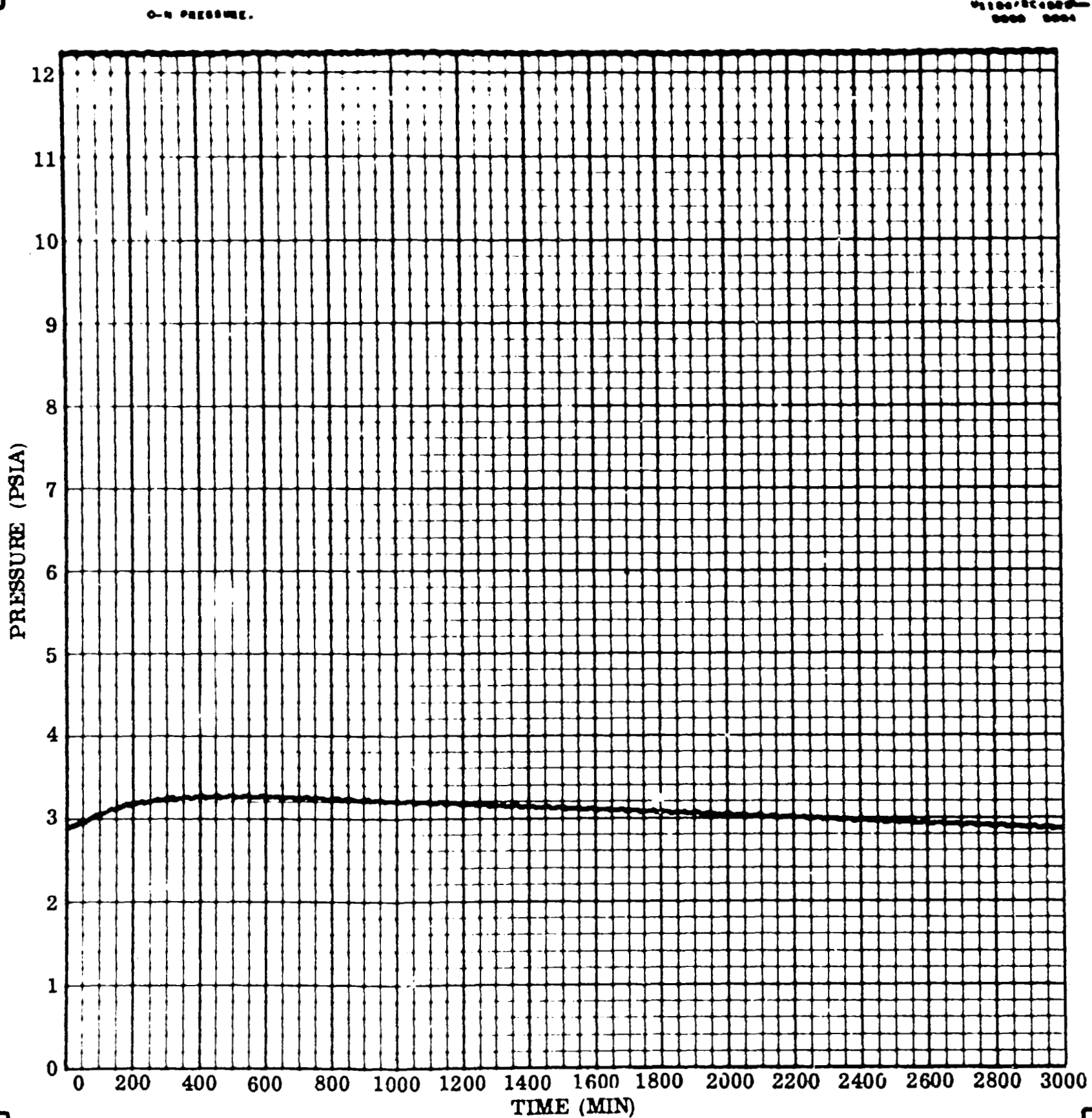


Fig. 2-41 Effect of Increased Hydrazine Feed Rate

FOLDOUT FRAME 4

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Test 3 Conditions. Several runs were made simulating the conditions of Test 3. Because many of the system parameters varied with time, average values of the parameters were used in the computer runs. The averaging process introduces small errors in that it assumes a linear relationship between current and rate constant. The best match that was achieved in three runs is shown in Fig. 2-42. The envelope and characteristic of the  $PO_2$  control are in good agreement with the Test 3 data (cycles 1, 2, 3, 5, and 6 of Test 3 are plotted).

In order to achieve agreement on mass balance, the computer case input for  $\dot{m}_{T_{out}}$  had to be decreased by approximately 10 percent. This is indicative of a combination of inaccuracies in metabolic/leak and nitrogen makeup instrumentation. This problem area was discussed previously in Section 2.3 under system analysis.

#### 2.4.2 Chaining

The versatility of the computer routine was increased by adding a chaining capability. This makes it possible in a single run to handle a sequence of step changes in case inputs. In effect, step changes in inputs during the run recycle the time base to make the conditions of  $P_T$ ,  $PO_2$ , and  $C$  at time =  $t$  become the initial conditions for the next time interval.

A computer run was made simulating the step changes of a portion of Test 4. The results are shown in Fig. 2-43. As with the simulation of Test 3, the envelope and characteristics of  $PO_2$  control are in good agreement, and the differences in cycle rate are attributed to errors introduced by the averaging process. The computer case input values for this run are given in Table 2-14. The case inputs are described in Ref. 1. The points in the run where the step changes were made are indicated in Fig. 2-43.

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### 2.4.3 Space Station Orbital Simulation

As part of the preliminary design of a 12-man  $O_2/N_2$  system for space station use, the computer routine was used to predict the control characteristics. A 94-min orbit with 58 min of daylight and 36 min of darkness was used. The high/low current mode on the sunlight side was set at nominal  $\pm 10$  percent and on the dark side at 50 percent of nominal. Fifty-three orbits are shown in Fig. 2-44. The long time constant shown for the  $PO_2$  control cycle indicates a need for the development of a more responsive  $PO_2$  control technique.

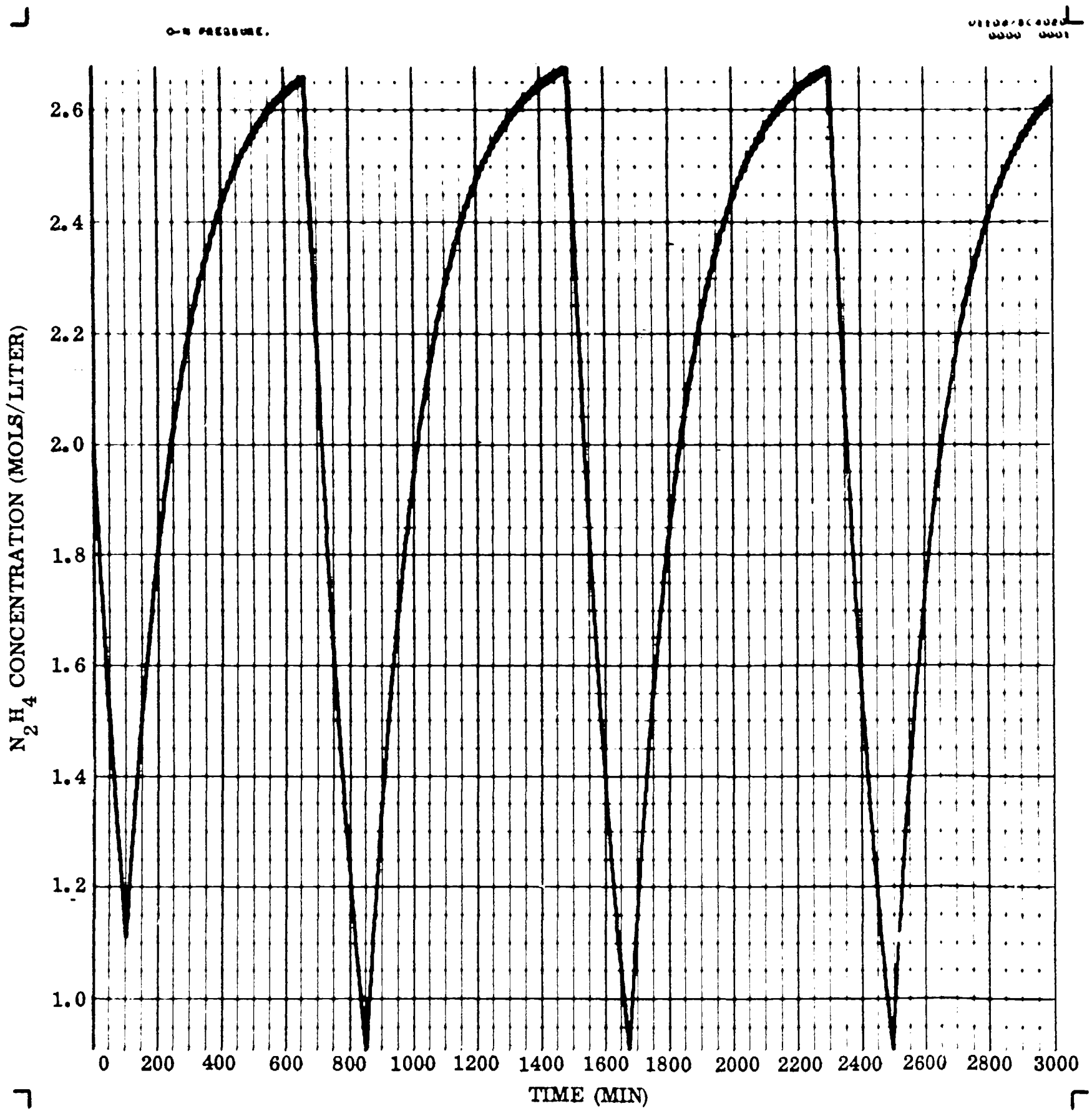
Table 2-14

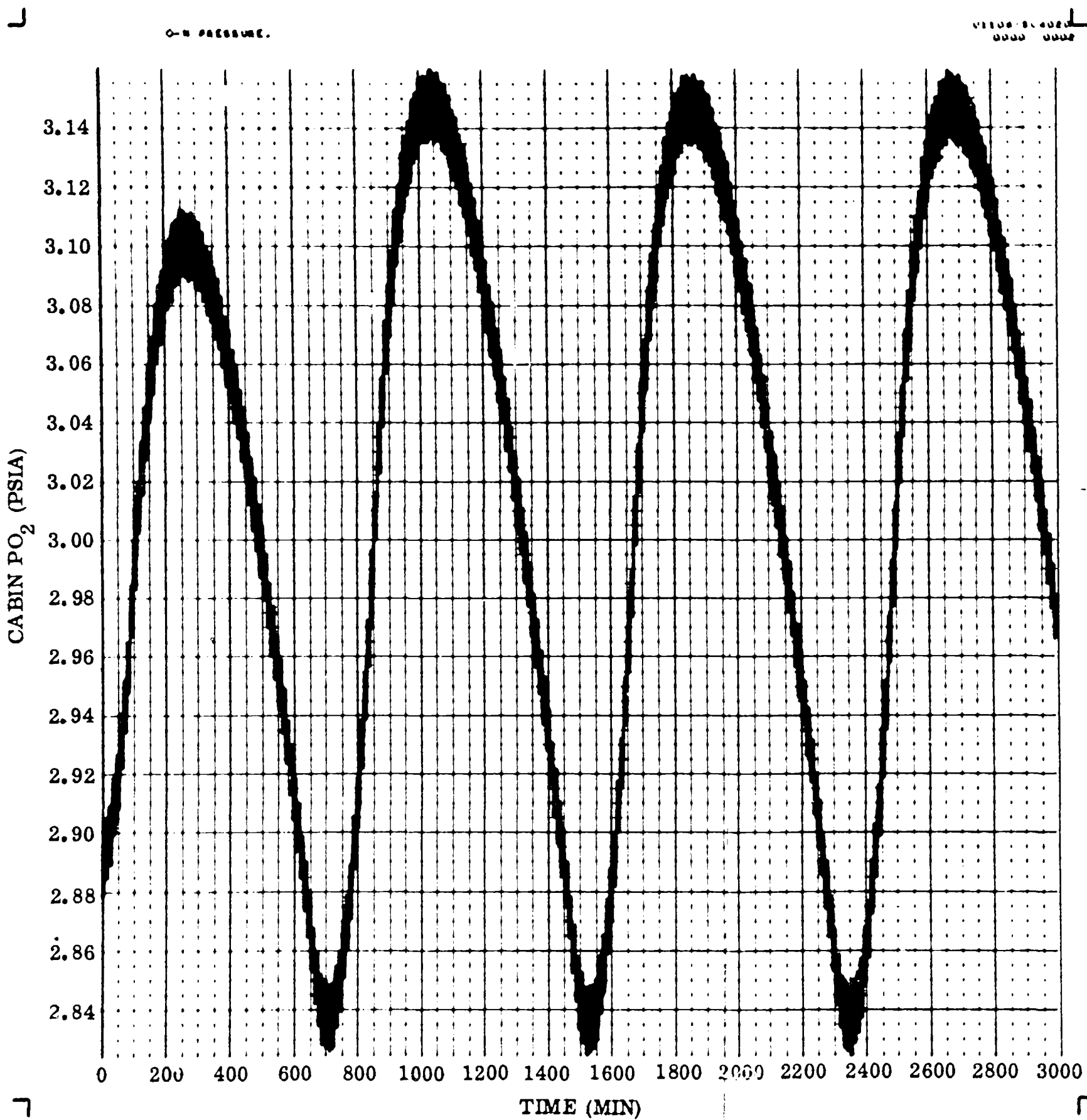
COMPUTER RUN 2-45 -- CASE INPUTS

Run No.	$T_0$	$T_{Max}$	$PO_2$ I	$P_{tot}$ I	$DMO_2$ Out	$DM_{tot}$ Out	$C_0$	$X_2$
2-45a	0	1,020	2.82	12.26	0.55	0.75	1.48	$1.4 \times 10^{-3}$
2-45b	1,020	1,110	-	-	0.31	0.51	-	$1.37 \times 10^{-3}$
2-45c	1,110	2,040	-	-	0.27	0.53	-	$1.37 \times 10^{-3}$
2-45d	2,040	2,325	-	-	0.37	0.68	-	$1.15 \times 10^{-3}$

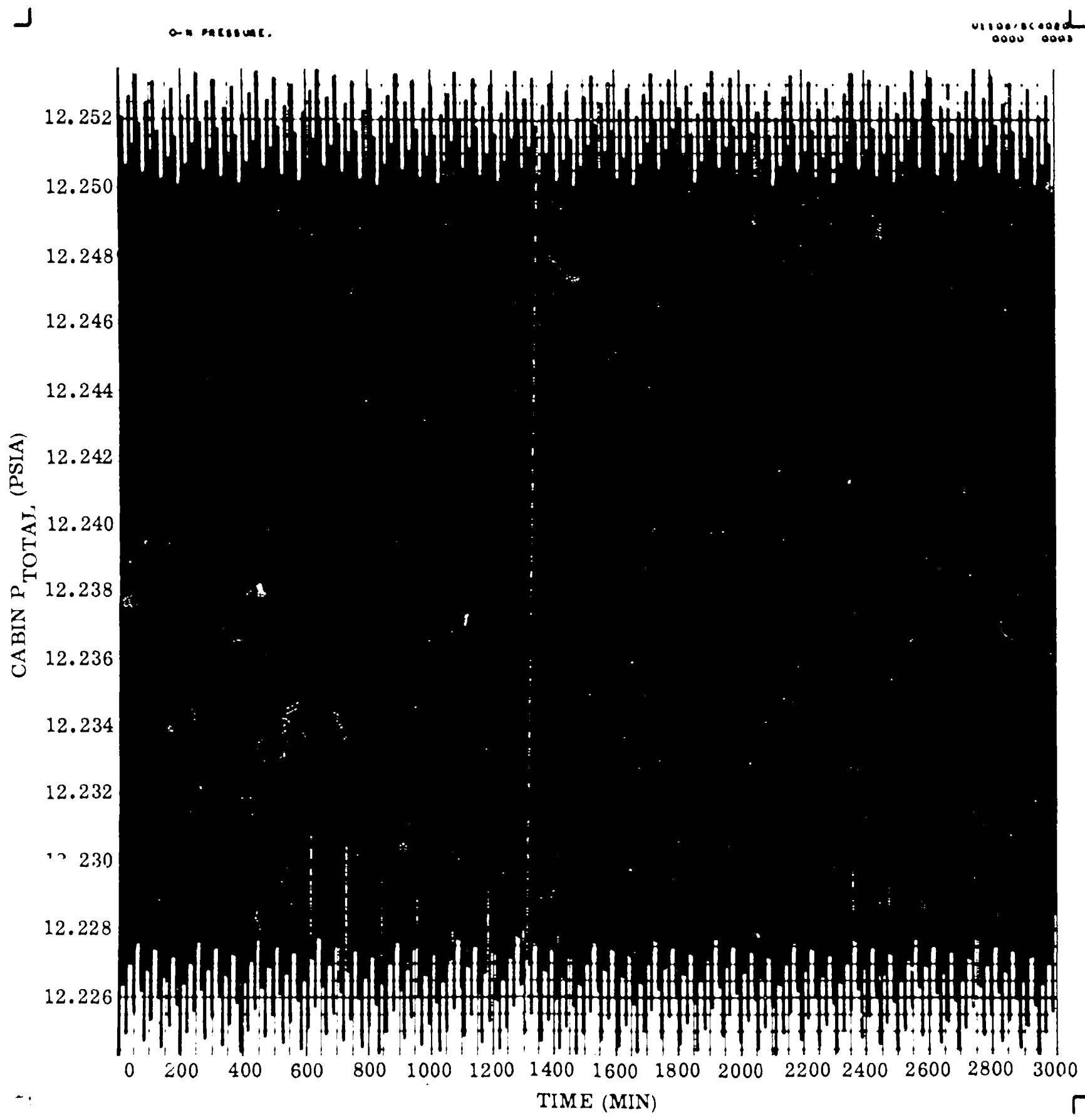
  

Run No.	$Y_1$	$Y_2$	$M_1$	$M_2$
2-45a	152	76	$4.35 \times 10^{-4}$	$3.66 \times 10^{-4}$
2-45b	153	72	$4.36 \times 10^{-4}$	$3.61 \times 10^{-4}$
2-45c	150	77	$4.32 \times 10^{-4}$	$3.66 \times 10^{-4}$
2-45d	153	74	$4.36 \times 10^{-4}$	$3.64 \times 10^{-4}$

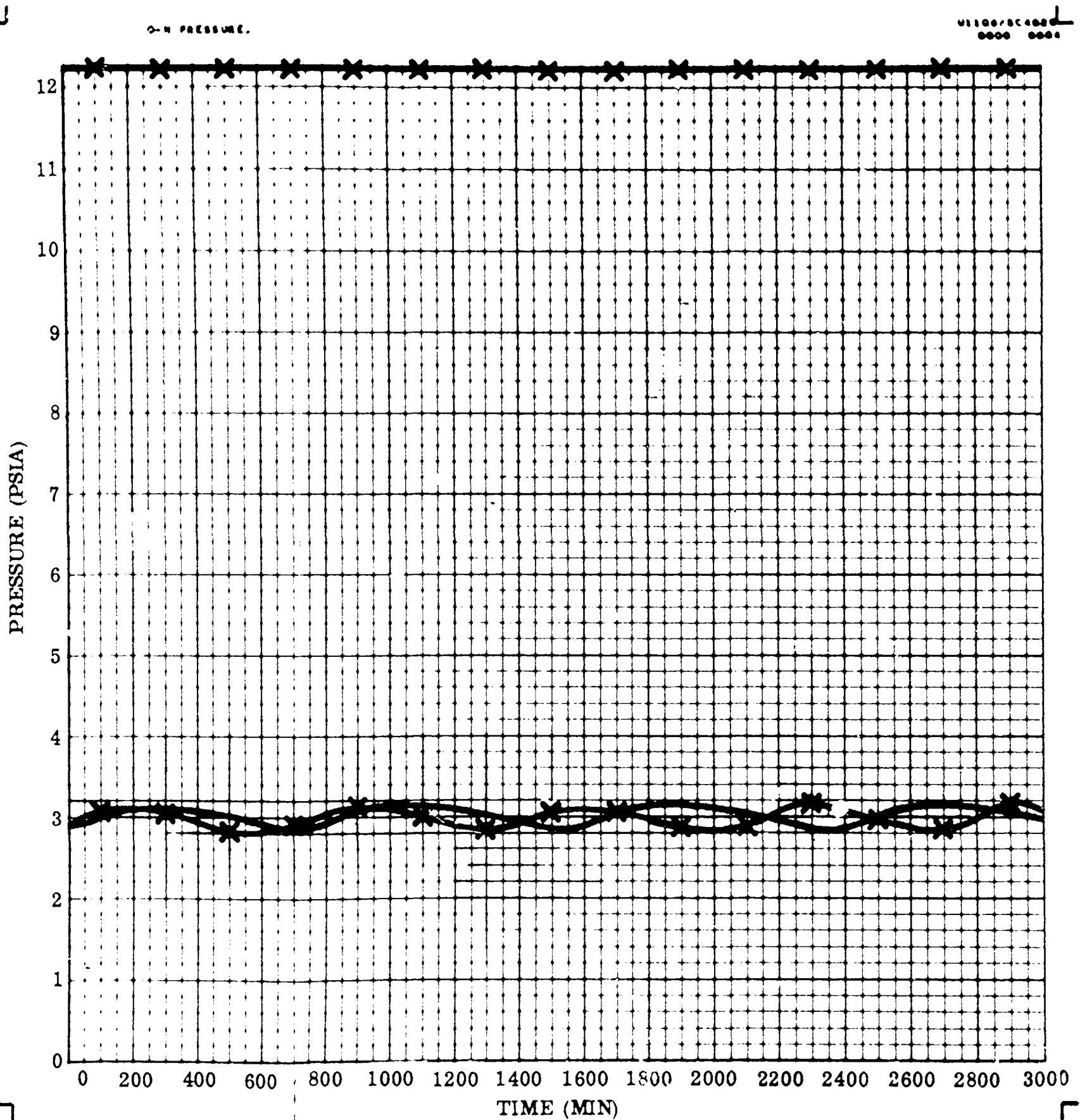




FOLDOUT FRAME 2



EOLDOUT FRAME 3



FOLDOUT FRAME 4

Fig. 2-42 Test 3 Simulation

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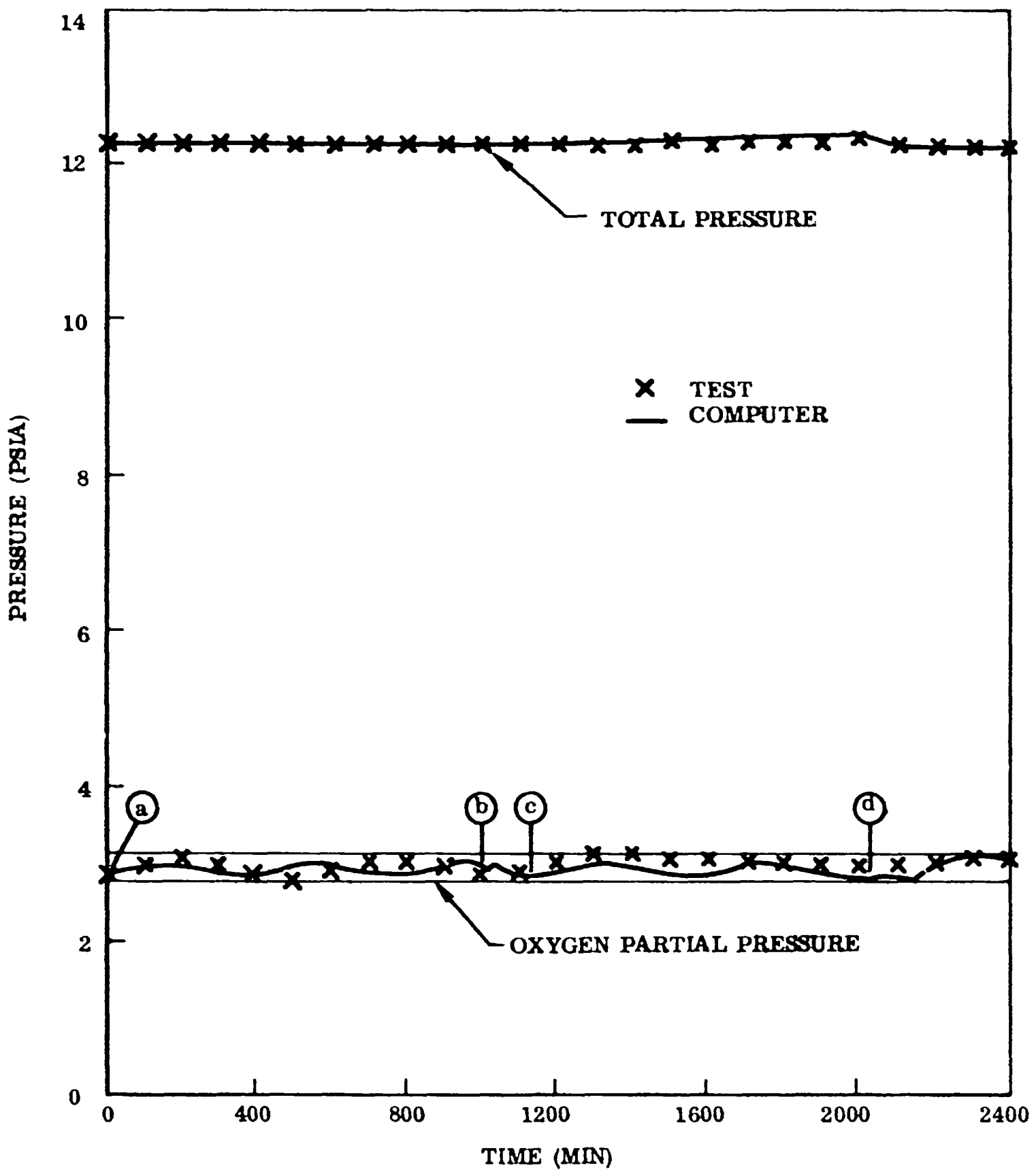
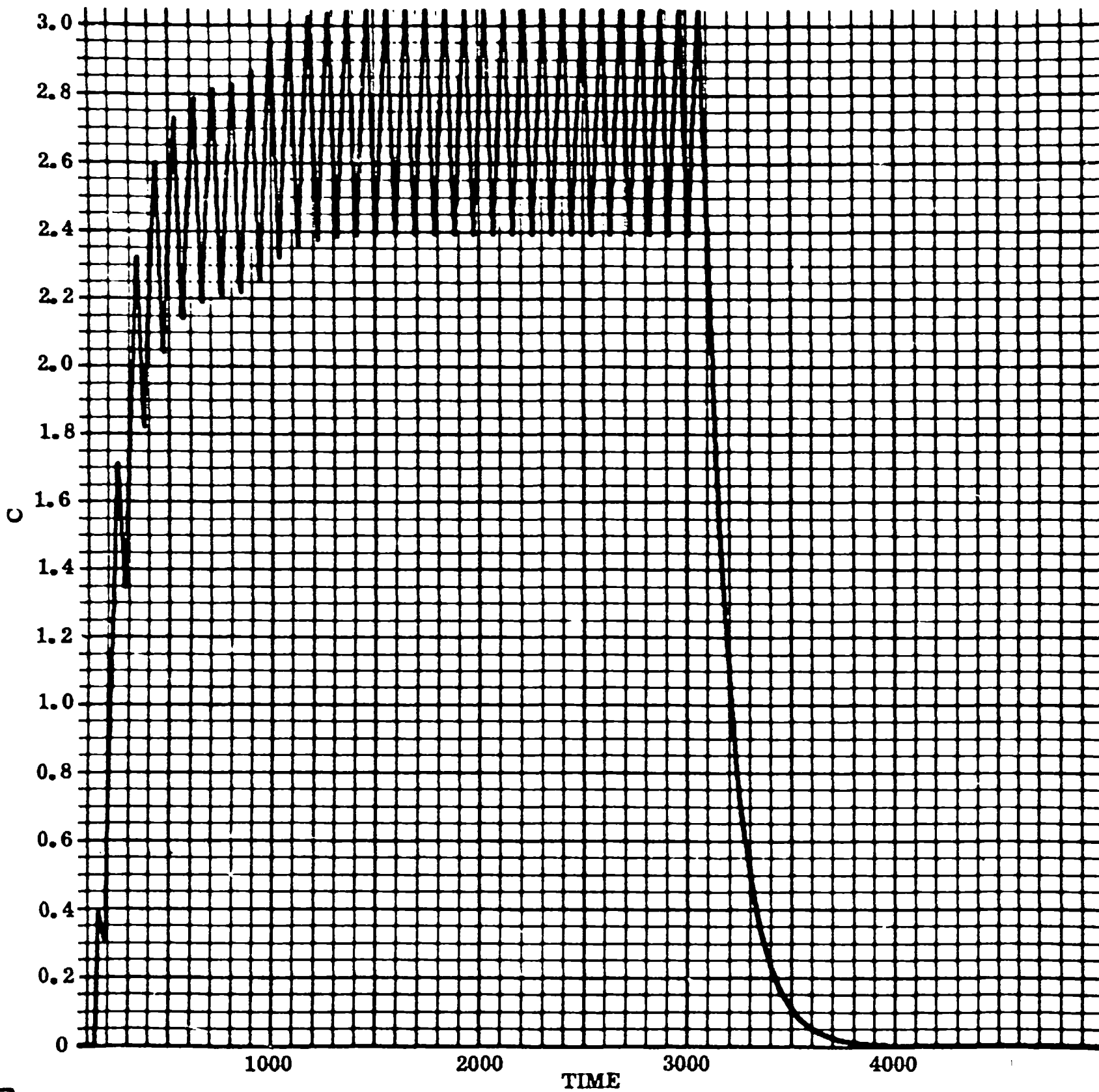


Fig. 2-43 Test 4 Simulation



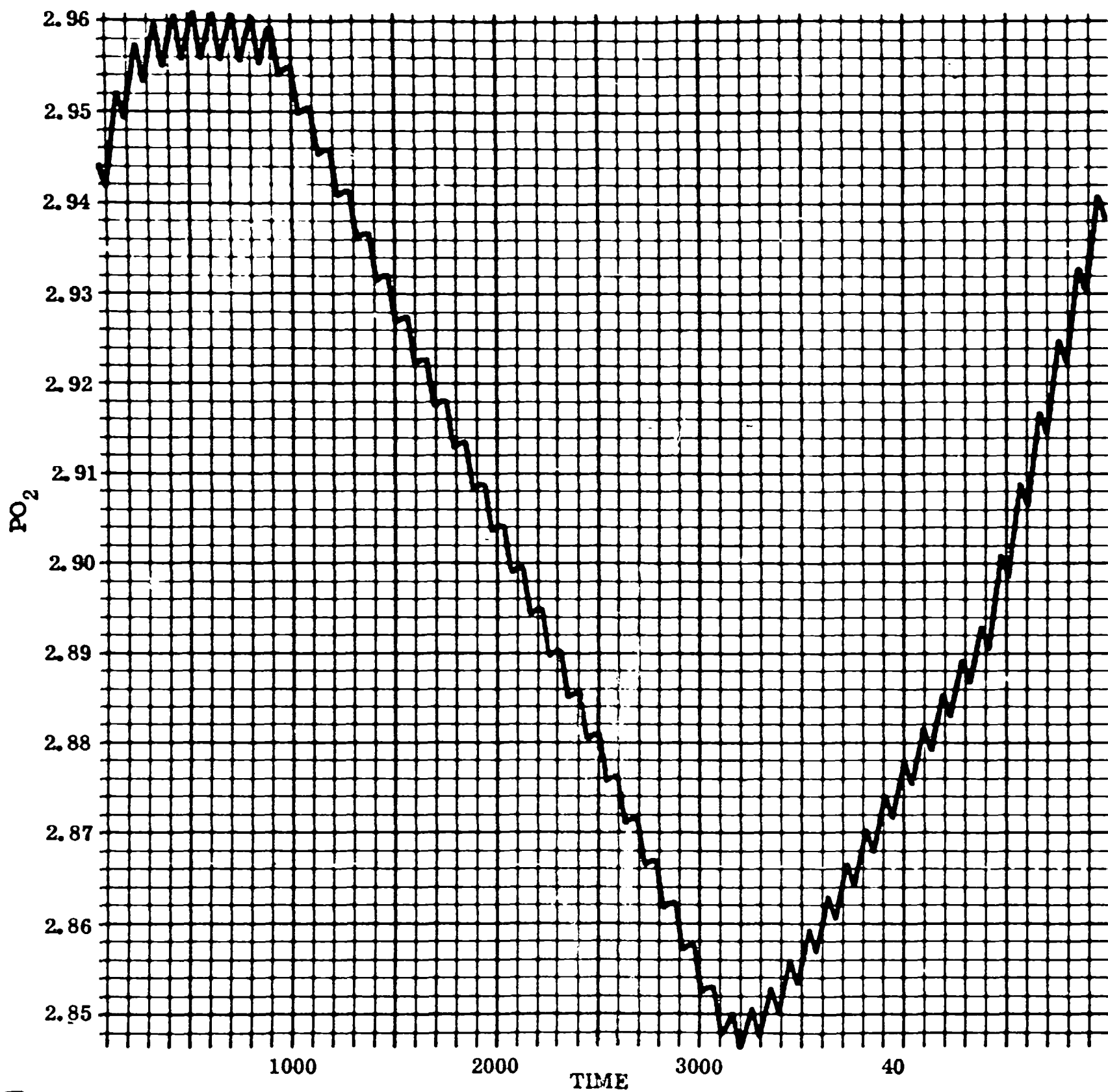
O-N PRESSURE

0000 0000



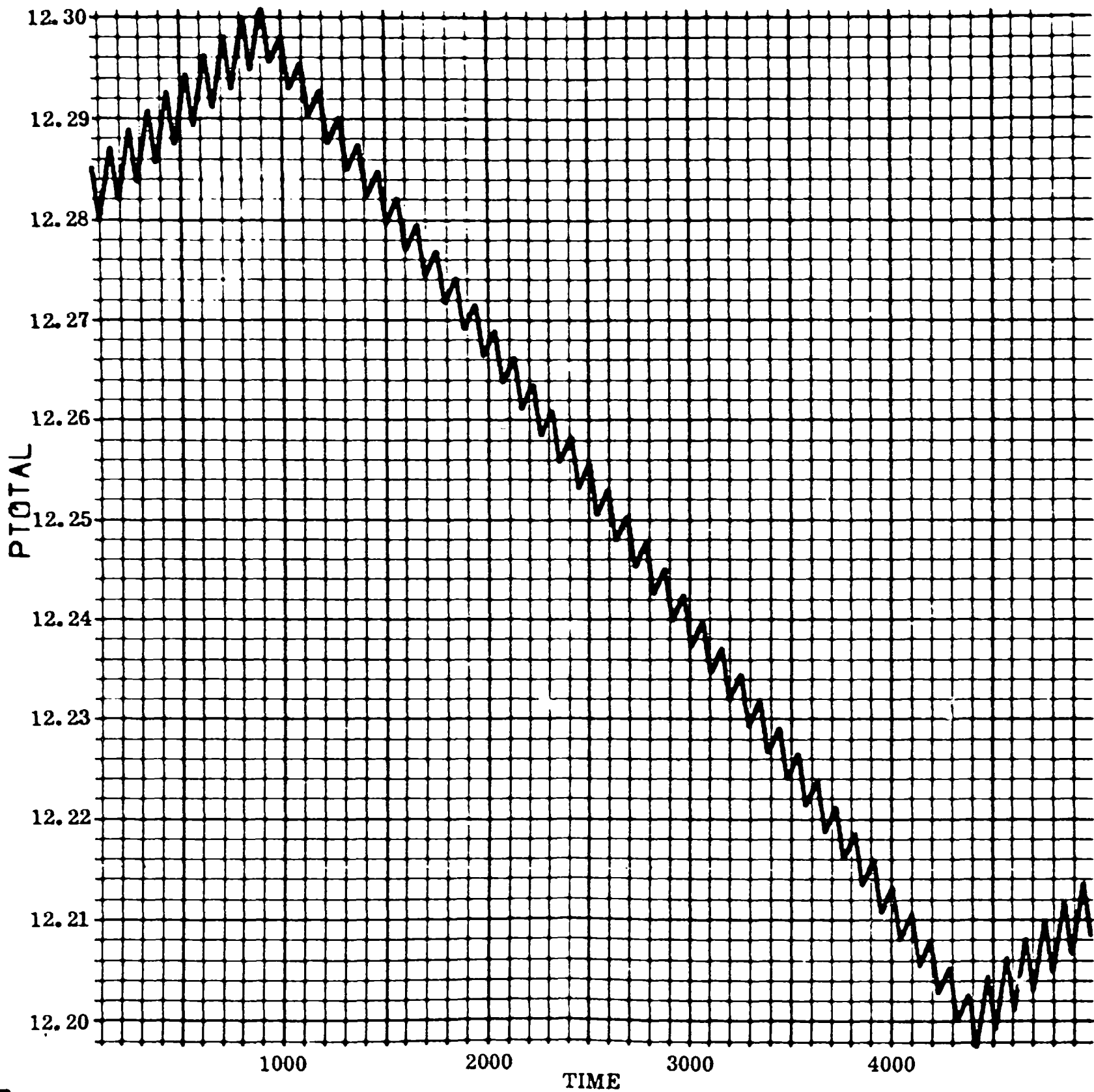
O-N PRESSURE

US100/SC4020  
0000 0000



O-M PRESSURE.

U3108/SC4020  
0000 0007



PRESSURE (PSIA)

O-N PRESSURE

P<sub>TOTAL</sub>

PRESSURE (PSIA)

CABIN PO<sub>2</sub>

1000

2000

3000

4000

TIME

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FOLDOUT FRAME 4

Fig. 2-44 Space Station Orbital Simulation

2-103



## 2.5 PRELIMINARY SYSTEM DESIGN

The preliminary design of a full-scale, 12-man oxygen/nitrogen generation system was conducted under this program utilizing the experimental data described in this report as a basis.

The preliminary design package included performance specifications, preliminary design layout drawings, interfaces and packaging, and supporting system analyses. A summary description of the preliminary system design and performance specification is presented in the following paragraphs.

### 2.5.1 PD System Description

In developing the design approach for the hydrazine water electrolysis system, the first consideration was an appropriate maintenance concept. Maintenance analysis revealed that the best approach was to avoid breaking into the electrolyte circuit to perform any maintenance tasks. This approach eliminates the necessity for handling the electrolyte in zero gravity and can be implemented with no significant weight penalty. The maintenance concept that was evolved consists of providing individual, self-contained hydraulic assemblies with no electrolyte connections; the electrolyte flow circuit is completely internally manifolded. These assemblies, or modules, are completely sealed to eliminate electrolyte leakage.

A reliability analysis was conducted to determine the optimum size and number of these individual modules. Various sizes and, hence, various numbers of modules were considered, ranging from one to nine active modules required for the 12-man capacity. Each module was first optimized in terms of redundant components to a point where no significant increase in reliability could be achieved by adding additional redundancy. The total number of modules, active plus spares, required to achieve a reliability goal of 0.9980 was then determined. These various systems were examined to determine which system required the least total weight.

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The analysis showed that a system employing nine active modules and five spare modules has the lowest weight. The approach to packaging these modules in a swing-out cabinet was to install 12 modules - nine active plus three installed spares - as shown in the Frontispiece of this report. The modules are mounted on rails to allow for easy installation and removal. Disconnects are provided in the rear of each module for the oxygen/nitrogen and hydrogen effluent gases, supply water and hydrazine, and the nitrogen required for pressurization. With this design, no electrolyte fittings need to be disconnected to replace a module. The cooling required for each module is provided by a conduction cold plate that interfaces between the module and the cabinet.

The swing-out cabinet also contains individual replaceable electronic assemblies for each module, water conditioning equipment, and a central status panel so that data supplied to the onboard computer system can be manually monitored.

The complete system to support 12 men for 180 days at the nominal metabolic and leakage rate with a reliability of 0.9980 requires a maximum of 7,830 watts of power and has a total fixed weight of 862 lb, including spares. The weight of an individual replaceable module is 46.6 lb.

#### 2.5.2 Summary System Specification

A summary of important performance parameters is presented in Table 2-15. The data presented include input power, waste-heat rejection rates, and inlet and outlet flow rates and pressures. Flow rates so noted represent the day-night average. Included are data for the nominal and the maximum cabin leak rates.

Table 2-15

## SUMMARY SYSTEM PERFORMANCE SPECIFICATION

Aspect	Nominal	Maximum
Oxygen generation rate (lb/day) <sup>(a)</sup>	26.1	33.2
Nitrogen generation rate (lb/day) <sup>(a)</sup>	8.0	22.6
Hydrogen generation rate (lb/day) <sup>(a)</sup>	4.75	7.59
Water consumption (lb/day) <sup>(a)</sup>	29.3	37.3
Hydrazine consumption (lb/day) <sup>(a)</sup>	9.72	27.5
Power consumption		
Day (unregulated 40 to 60 Vdc)(W)	7,830	13,300
Night (regulated 28 Vdc)(W)	3,180	5,160
Heat rejection		
Day (Btu/hr)	13,700	23,900
Night (Btu/hr)	4,720	7,960
N <sub>2</sub> supply (lb/day)	0.07	0.09
Output pressure, H <sub>2</sub> and O <sub>2</sub> /N <sub>2</sub> , variable (psia)	1 to 12	1 to 12
Operating temperature (° F)	75	75
Number of active modules	9	12
Total system weight, with spares (lb)	862	862
Weight of replaceable module (lb)	46.6	46.6

<sup>(a)</sup>Day-night average

### Section 3 CONCLUSIONS

#### 3.1 ELECTRODE DEVELOPMENT

Experimental electrodes were developed with performance comparable to the commercial material used in previous programs. Their suitability for use in a hydrazine electrolysis system was demonstrated by the performance of three of these electrodes as anodes in the one-man model  $O_2/N_2$  system testing. Exploratory testing of reduced-catalyst cathodes also indicated an approach to improvement in hydrazine reaction efficiency.

#### 3.2 ONE-MAN MODEL $O_2/N_2$ SYSTEM

The one-man model  $O_2/N_2$  generator integrated with a cabin and metabolic/leak simulator provided an excellent testbed for evaluating the hydrazine water electrolysis approach to a noncryogenic  $O_2/N_2$  generation system. Significant data concerning component performance, system control techniques, and cabin atmosphere control characteristics were obtained. Adequate control of the cabin simulator total pressure and oxygen partial pressure were demonstrated.

#### 3.3 COMPUTER MODEL

The computer routine model, developed under Contract NAS1-7706, was revised and updated based on the system test data. Reasonable agreement was obtained between the test and computer prediction of space cabin atmosphere control. The revised computer routine predicts adequate control of a full-scale 12-man space station cabin.

### 3.4 PRELIMINARY SYSTEM DESIGN

A preliminary design effort with emphasis on maintainability, reliability, a sound basis of experimental data and proven concepts resulted in the preliminary design and specification of a 12-man  $O_2/N_2$  generation system suitable for space station use. The complete system to support 12-men for 180 days at the nominal metabolic and leakage rate with a reliability of 0.9980 requires a maximum of 7,830 watts of power and has a total fixed weight of 862 lb, including spares. The weight of an individual replaceable module is 46.6 lb. The system is capable of handling a maximum cabin leakage of 33.7 lb/day.

Section 4  
REFERENCES

1. Lockheed Missiles & Space Company, Utilization Manual Nitrogen Generation Control Analysis, by K. G. Barany, (OXNIP) BRCWOO, Sunnyvale, Calif., Mar 1970
2. -----, Preliminary Design of a Space Station Electrolytic Oxygen-Nitrogen Generator, by B. M. Greenough, LMSC-A977498, Sunnyvale, Calif., 5 Mar 1971
3. W. J. Conner, B. M. Greenough, and G. M. Cook, "Design and Development of a Water Vapor Electrolysis Unit," NASA CR 607, Sep 1966
4. R. G. Haldeman, "Electrode-Matrix Materials," reprinted from 21st Annual Proceedings Power Sources Conference, May 1967
5. B. M. Greenough, "The Development and Preliminary Design of an Oxygen-Nitrogen Generation System," NASA CR 66940 Jun 1970
6. Andrith and Ogg, The Chemistry of Hydrazine, John Wiley & Sons, New York, 1951
7. B. M. Greenough and T. M. Olcott, "A Spacecraft Electrolytic Oxygen-Nitrogen Generation System," presented at the Space Technology and Heat Transfer Conference, ASME, Jun 1970

### LIBRARY CARD ABSTRACT

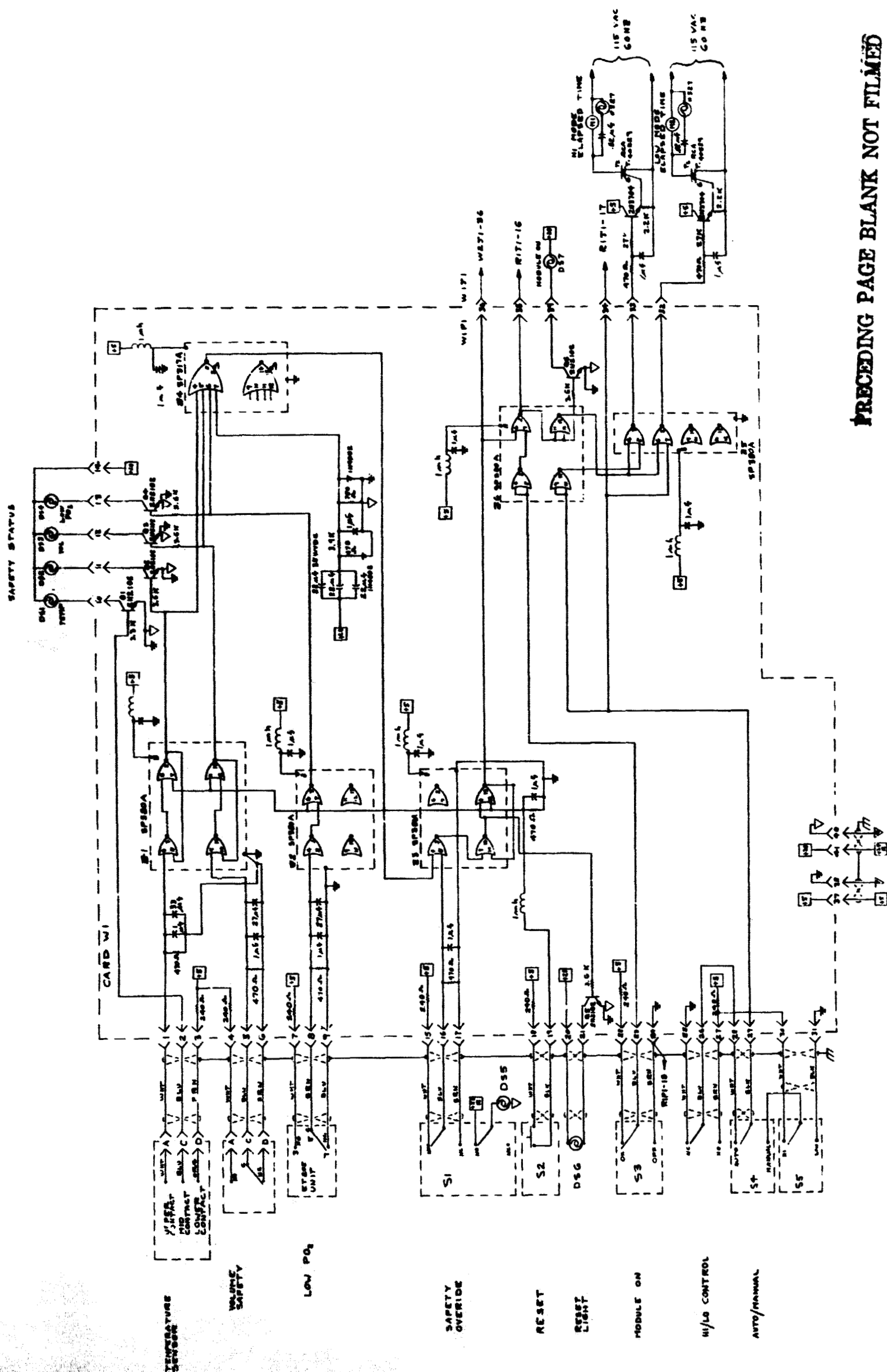
Research was conducted in the development of a noncryogenic technique of hydrazine/water electrolysis for generating oxygen and nitrogen for use in long-duration manned space missions. A breadboard one-man model  $O_2/N_2$  generation system was integrated with a cabin and metabolic/leak simulator. Automatic control of the cabin total pressure and cabin oxygen partial pressure was demonstrated for various metabolic loads with a fixed cabin leakage. A total of four one-week tests were run. Experimental electrodes were tested in the system in addition to some 5,000 hr of cumulative single-cell tests. A computer routine model of the  $O_2/N_2$  generation system was revised and verified with experimental test data. A preliminary design and specification of a full-scale 12-man  $O_2/N_2$  system was completed yielding a system capable of handling up to 33.7 lb/day cabin leakage and a 12-man metabolic load. Total system weight, including spares, is 862 lb. The system requires a maximum of 7,830 watts under nominal conditions and has redundancy and spares for a reliability of 0.9980 for a 180-day mission.

**Appendix A**  
**ONE-MAN MODEL SYSTEM CIRCUIT DIAGRAMS**

Circuit diagrams of the control logic, power conditioning, and voltage/current monitors used in the one-man model  $O_2/N_2$  system are presented in this Appendix.







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Fig. A-1 Control Logic Card 1

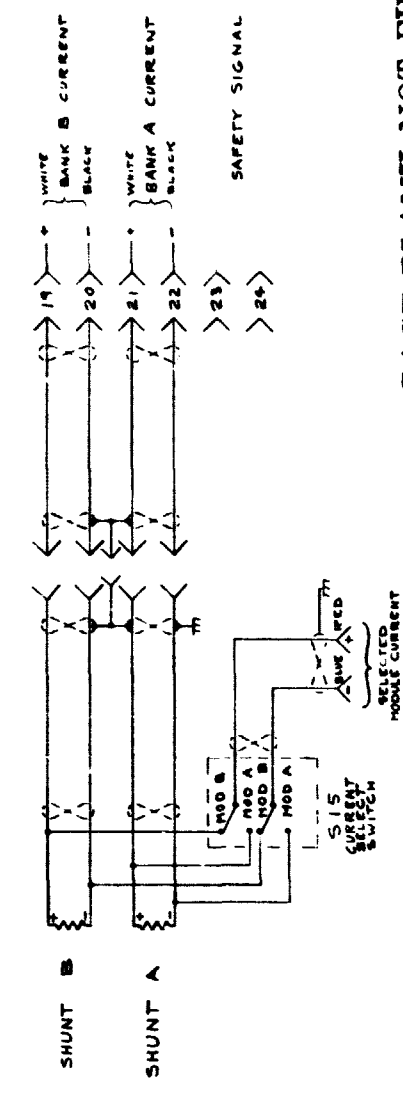
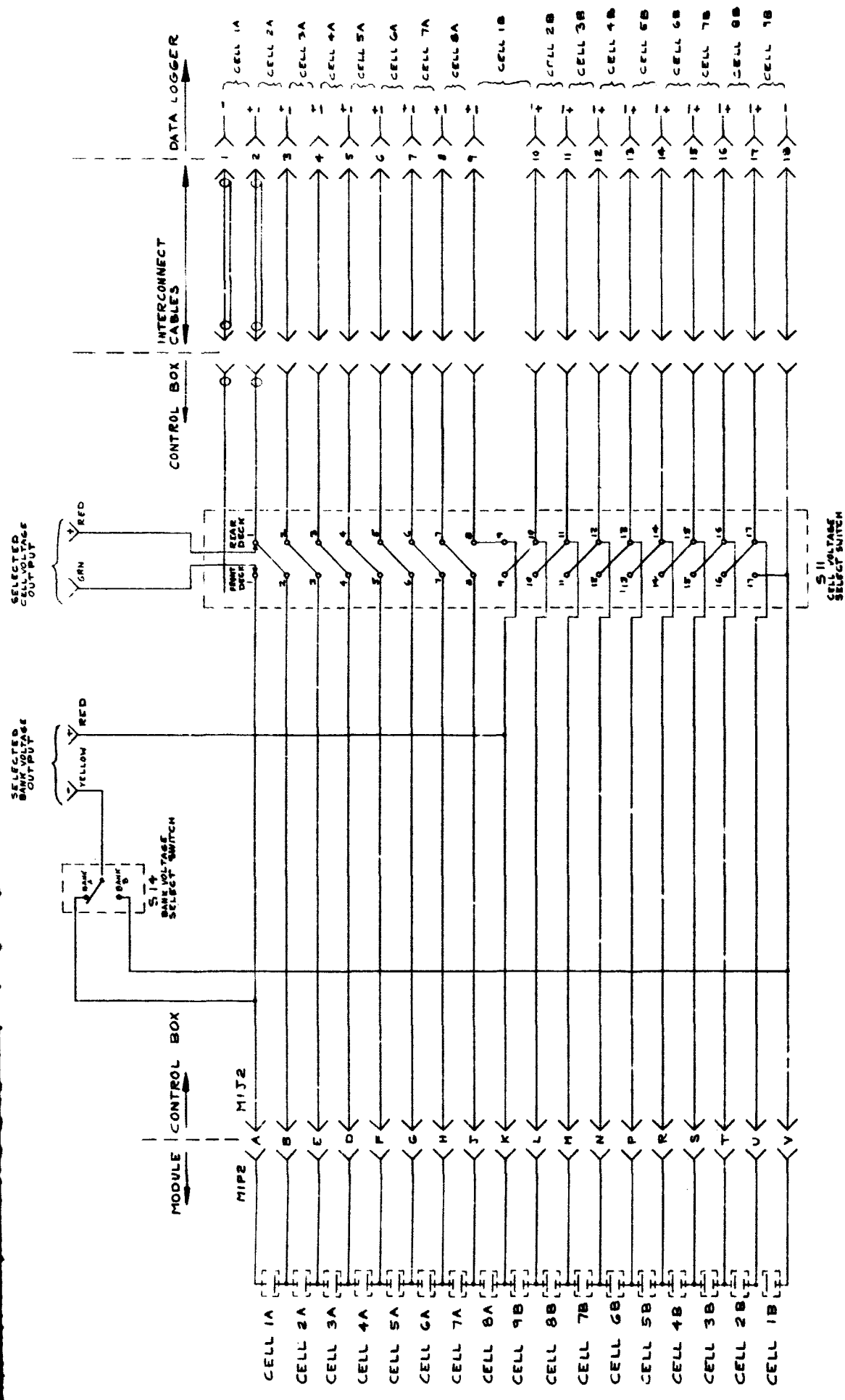












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Fig. A-4 Voltage/Current Monitor  
A-9

## Appendix B

### OPERATION INSTRUCTIONS FOR THE ONE-MAN MODEL O<sub>2</sub>/N<sub>2</sub> SYSTEM

#### A. Manual Shutdown

1. Emergency shutdown: Disconnect labelled power supply cords behind monitoring consoles.
2. Normal shutdown:
  - a. Turn off N<sub>2</sub>H<sub>4</sub> feed switch.
  - b. Turn off H<sub>2</sub>O feed switch.
  - c. Turn on N<sub>2</sub> purge switch.
  - d. Switch to manual, low current mode on electrolysis module control panel.
  - e. Turn off module power switch.
  - f. Turn off dia-pump switch.
  - g. Turn off KOH pump switch.
  - h. Turn off power supplies in this order:
    - (1) 28 Vdc (PS 3)
    - (2) 28 Vdc (PS 2)
    - (3) 5 Vdc (PS 1)
  - i. Press POWER OFF button.
  - j. Pin reservoir.
  - k. Check N<sub>2</sub> purge flow rate and adjust.

#### B. Manual Start-up

1. Switch positions:
  - a. Module power - off
  - b. Module mode - manual, low
  - c. Safety override - on
  - d. N<sub>2</sub>H<sub>4</sub> feed - off
  - e. N<sub>2</sub>O feed - off
  - f. N<sub>2</sub> purge - on
  - g. KOH pump - off
  - h. Dia-pump - off



2. Press POWER ON button.
3. Turn on power supplies in this order:
  - a. 5 Vdc (PS 1)
  - b. 28 Vdc (PS 2)
  - c. 28 Vdc (PS 3)
4. Press RESET button.
5. Turn on KOH pump switch: Adjust variac to 70 or flow of 5 units on flowmeter C. Check to see that bubble separator gas line is not drawing in  $H_2O$ ; adjust flow downward with flow control valve if there is suction pressure or bubble separator.
6. Turn on  $H_2O$  feed switch.
7. Turn on module power switch; check that low currents (modules A and B) read  $\sim 11.2$  mV.
8. Switch to manual high current; check that currents (modules A and B) read  $\sim 33.8$  mV.
9. Turn on dia-pump switch. When  $P_{Total}$  reaches 12.3 psia ( $\sim 61.5$  on recorder), do the following:
  - a. Switch to automatic current.
  - b. Turn off  $N_2$  purge.
  - c. Turn off safety override.
  - d. Set metabolic leak to \_\_\_\_\_ cc/min. (\_\_\_\_\_ SS).
  - e. Set  $N_2$  makeup to \_\_\_\_\_ cc/min. (\_\_\_\_\_ SS).
  - f. Turn on  $N_2H_4$  feed to \_\_\_\_\_ cc/min. (\_\_\_\_\_ SS).
  - g. Set pressure regulator to 1 psig.

C. Start-up From Automatic Shutdown

1. Switch positions:
  - a. Module power – off
  - b. Module mode – manual, low
  - c. Safety override – on
  - d.  $N_2H_4$  feed – off
  - e.  $H_2O$  feed – off
  - f.  $N_2$  purge – on

- g. KOH pump – off, variac 0
- h. Dia-pump – off
- 2. Press RESET button.
- 3. Turn on KOH pump switch; adjust flow with variac (~70 V).
- 4. Turn on H<sub>2</sub>O feed switch.
- 5. Turn on module power switch; check low currents (~11.2 mV), then check high currents (~33.8 mV).
- 6. Turn on dia-pump; adjust metabolic leak and N<sub>2</sub> makeup.
- 7. Turn off N<sub>2</sub> purge switch.
- 8. Turn on N<sub>2</sub>H<sub>4</sub> feed.

#### D. Calibration

- 1. F3 O<sub>2</sub> Analyzer
  - a. Open cal vent valve.
  - b. Set cal vent 3-way valve in vent position.
  - c. Set cal loop 3-way valve in cal position.
  - d. Set O<sub>2</sub>/N<sub>2</sub> 3-way valve in N<sub>2</sub> position.
  - e. Adjust flow through cal loop flowmeter to ~50 cc/min.
  - f. Allow ~10 to 15 minutes for purging of F3; then adjust F3 zero knob to read 0 percent.
  - g. Set O<sub>2</sub>/N<sub>2</sub> 3-way valve in O<sub>2</sub> position.
  - h. Allow ~10 to 15 minutes for purging of F3; then adjust F3 span knob to read 100 percent.
  - i. Return 3-way valves to normal cal loop positions:
    - (1) Cal loop 3-way valve in loop position
    - (2) Cal vent 3-way valve in loop position
    - (3) O<sub>2</sub>/N<sub>2</sub> 3-way valve in N<sub>2</sub> position.
- 2. PO<sub>2</sub> Control Sensor and Data Sensor
  - a. Turn off N<sub>2</sub>H<sub>4</sub> feed switch.
  - b. Turn on sensor bypass valve and turn off sensor loop valves.
  - c. Disconnect sensor cables and remove sensors from canister, then reconnect sensor cables.

- d. Turn control amplifier to zero position and adjust zero reading; turn to 250 mm position and set calibration to give meter reading of 118 mm. Repeat same procedure for data sensor; also check that PO<sub>2</sub> recorder reads correctly (the zero can be adjusted on the recorder).
  - e. Replace sensors in canister; turn on sensor valves and turn off bypass valve.
  - f. Turn on N<sub>2</sub>H<sub>4</sub> feed switch.
  - g. If sensor is reading above 154 mm, note reading and turn calibration knob below 154 mm, then back to reading noted.
3. PO<sub>2</sub> Safety sensor
- a. Turn on safety override switch.
  - b. Turn on sensor bypass valve and turn off sensor loop valves.
  - c. Remove sensor from canister.
  - d. Turn amplifier to zero position and adjust zero reading; turn to 1,000-mm scale and set reading with calibration knob to 160 mm.
  - e. Replace sensor in canister; turn on sensor loop valves and close bypass valve.
  - f. Turn off safety override switch.

#### E. Current Adjustment

- 1. Adjust low current as follows:
  - a. Switch on manual low current.
  - b. Connect voltmeter to module current readout.
  - c. Put module current-selector switch on "A" and adjust "low A" potentiometer on Electronics Card 1 to read ~11.2 mV.
  - d. Switch to "B" and adjust "low B" potentiometer on Card 1 to read ~11.2 mV.
- 2. Adjust high current as follows:
  - a. Switch to high current on the electrolysis module.
  - b. Turn current-selector switch to "A" and adjust "A high" potentiometer on Electronics Card 1 to ~33.3 mV.
  - c. Switch to "B" and repeat.

F. Filling  $\text{H}_2\text{O}$  and  $\text{N}_2\text{H}_4$  Tanks

1. Fill  $\text{H}_2\text{O}$  tank as follows:

- a. Turn off  $\text{H}_2\text{O}$  feed switch.
- b. Close off tank pressurization valve to isolate  $\text{H}_2\text{O}$  tank from  $\text{N}_2$  supply.
- c. Open  $\text{H}_2\text{O}$  vent valve to depressurize  $\text{H}_2\text{O}$  tank.
- d. Drain  $\text{H}_2\text{O}$  tank through  $\text{H}_2\text{O}$  fill line and measure volume of  $\text{H}_2\text{O}$ .
- e. Connect  $\text{H}_2\text{O}$  fill bottle to  $\text{H}_2\text{O}$  fill line.
- f. Open  $\text{H}_2\text{O}$  fill valve.
- g. Pump in measured amount of distilled  $\text{H}_2\text{O}$ .
- h. Close  $\text{H}_2\text{O}$  fill valve.
- i. Close  $\text{H}_2\text{O}$  vent valve.
- j. Open tank pressurization valve to pressurize  $\text{H}_2\text{O}$  tank.
- k. Turn on  $\text{H}_2\text{O}$  feed switch.

2. Fill  $\text{N}_2\text{H}_4$  tank as follows:

- a. Turn off  $\text{N}_2\text{H}_4$  feed switch.
- b. Open  $\text{N}_2\text{H}_4$  vent valve to depressurize  $\text{N}_2\text{H}_4$  tank. Keep  $\text{N}_2$  purge flowing over  $\text{N}_2\text{H}_4$  tank.
- c. Drain  $\text{N}_2\text{H}_4$  from  $\text{N}_2\text{H}_4$  tank through  $\text{N}_2\text{H}_4$  fill line, and measure volume.
- d. Connect  $\text{N}_2\text{H}_4$  fill bottle to  $\text{N}_2\text{H}_4$  fill line.
- e. Open  $\text{N}_2\text{H}_4$  fill valve.
- f. Pump in measured volume of  $\text{N}_2\text{H}_4$ , but do not pump in air. Measure remaining  $\text{N}_2\text{H}_4$  in fill bottle to determine amount of  $\text{N}_2\text{H}_4$  introduced into tank.
- g. Close  $\text{N}_2\text{H}_4$  fill valve.
- h. Close vent valve to pressurize  $\text{N}_2\text{H}_4$  tank.
- i. Turn on  $\text{N}_2\text{H}_4$  feed switch.

Appendix C  
BREADBOARD SYSTEM TEST DATA LOGS

TEST NO. 1

## SYSTEM STATUS

PAGE 2 OF 2

DATE	TIME	ELAPSED TIME (HRS)	HYDROGEN OUTPUT METER		POWER INPUT		PRESSURE (PSIG)			COLLANT TEMP (°F)	FLOW RATE			GAS ANALYSIS									
			HIGH VOLTS	LOW VOLTS	HIGH AMPS	LOW AMPS	MUD PI	PO2	LAL		NH4	KCH	WANTIN	HA	ETA	N2	N2-O2	CASIN	HL				
9-22	4:15	0	2192.2		28.5	6.8	40	5.0	135	2.2	6.1	5.9	6.0	5.5	575	2040	1690						
	5:00	~1	1483.1		28.5	6.0	39	4.0	137	2.2	6.1	5.9	6.58	5.5	575	2040	1690						
	8:05	4	1495.2		28.5	21.5	34	5.0	150	2.0	6.0	49	5.4	5.5	575	2040	1690						
	11:40	7:35	1502.6		28.6	5.5	23	3.1	166	2.1	6.0	53	3.8	—	575	2040	1690						
9-23	12:40	8:35	1511.5		28.5	5.8	26	4.2	170	2.1	6.0	50	5.3	—	575	2040	1690						
	10:31	9:30		28.6	21.2		27	4.2	187	2.1	5.8	50	5.0	—	575	2040	1690						
	12:32	10:30	1518.3		28.6	22.5	25	4.2	213	2.1	5.9	50	4.3	5	575	2040	1690						
	3:35	11:30	1521.8		28.7	23.0	23	4.2	233	2.1	5.9	49	5.0	5	575	2040	1690						
	4:30	12	1525.1		28.8	22.0	36	4.2	253	2.1	5.9	50	5.0	5	575	2040	1690						
	5:30	13	1528.0		28.7	23.0	33	4.2	273	2.1	5.8	50	5.2	0	575	2040	1690						
	6:30	14	1532.5		28.7	23.6	31	4.1	289	2.1	5.8	49	5.0	5	575	2040	1690						
	10:45	18.2	1546.5		28.7	2.5	52	4.1	40%	2.1	5.9	49	4.9	0	575	2040	1690						
	12:45	20.2	1548.5		28.5	19	42	4.6	31%	2.1	5.9	49	5.0	0	575	2040	1690						
	2:45	22.2	1564.0		28.5	19.5	33	5.5	170%	2.1	5.9	48	4.5	0	575	2040	1690						
	4:45	24.2	1574.1		28.5	19.5	31.5	4.2	151	2.1	5.9	47	3.7	5	575	2040	1690						
	6:45	26.2	1578.1		28.5	6.0	33.5	4.2	152	2.1	5.9	50	4.2	0	575	2040	1690						
	8:45	28.2	1583.1		28.5	5.9	32.0	4.0	137	2.1	5.9	52	4.0	0	575	2040	1690						
	10:45	30.2	1586.1		28.5	5.9	35.5	4.0	126	2.1	5.8	54	4.0	0	575	2040	1690						
9-24	12:35	32	1588.3		28.5	5.7	37.7	4.0	126	2.1	5.9	54	4.0	0	575	2040	1690						
	2:35	34	1591.0		28.5	5.2	35.0	4.0	127	2.1	5.9	56	4.0	0	575	2040	1690						
	4:35	36	1593.4		28.6	5.4	33.5	4.0	130	2.1	5.95	58	4.0	0	575	2040	1690						
	6:35	38	1595.5		28.6	5.7	36.0	4.0	—	2.1	6.0	55	4.0	0	575	2040	1690						
	10:45	42.2	SEE LOG																				
	10:50		1601.7			28.6	5.7	33	4.1	161	2.1	5.9	52	4.0	—	575	2040	1690					
	12:50	44	1609.5			28.5	20.0	28.5	5.7	34	4.0	157	2.1	5.9	16	575	2040	1690					
	1:30	45	1626.5			28.5	19.9	31	4.1	149	2.1	5.9	49	3.8	5	575	2040	1690					
	3:40	47.2										48	3.8	6	72	575	2040	1690					
	3:45																						
	4:45																						
	5:25	49																					





TEST NO. 1

MODULE STATUS

PAGE 1 OF 1

DATE	TIME	HIGH TIMER	LOW TIMER	MODE	MOD VOLTS		MOD AMPS		CELL VOLTAGE												TOTAL VOLTS	PERCENT	REMARKS				
					A	B	A	B	1A	2A	3A	4A	5A	6A	7A	8A	9A	10A	11A	12A							
9-22	4:30	9:59	7:43	L05	13.8	14.98	4.3	4.4	1.671	1.765	1.821	1.629	1.905	1.763	1.748	1.600	1.704	1.442	1.623	1.693	1.648	1.661	1.665	1.677	31.90(5)		
	5:30	10:06	8:06	H15	16.0	17.45	13.6	13.6	2.033	2.213	2.105	1.813	1.995	2.008	2.061	1.772	1.975	1.824	1.822	1.960	1.864	1.913	2.010	2.003	2.077	33.97	0.64 (6.42)
	7:00	11:26	8:06	H15	15.79	17.54	13.6	13.5	2.165	2.269	2.134	1.789	1.820	1.830	2.027	1.753	1.932	1.784	1.790	1.908	1.837	1.900	2.096	2.121	2.190		1.31 (8.00)
	8:30	12:40	8:06	H15	15.76	17.72	14.0	13.8	2.228	2.308	2.160	1.803	1.840	1.832	2.035	1.759	1.923	1.783	1.798	1.908	1.840	1.908	2.144	2.194	2.241	29.70	
	9:45	13:64	8:06	H15	15.81	17.59	13.4	13.5	2.215	2.281	2.145	1.795	1.820	1.807	1.997	1.748	1.906	1.761	1.769	1.894	1.829	1.893	2.132	2.182	2.220	23.90	12.7 (10.00)
	11:40	14:92	8:06	L05	11.17	11.16	13.44	14.98	1.71	1.80	1.70	1.62	1.66	1.64	1.73	1.60	1.62	1.61	1.60	1.60	1.64	1.66	1.68	1.69	1.73	23.70	18.5 (10.00)
9-23	12:00																										
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TEST NO. 1

## MODULE STATUS

PAGE 2 OF

DATE	TIME	HIGH TIMER	LOW TIMER	MODE	MOD VOLTS		MOD AMPS		CELL VOLTAGE												EFFICIENCY %	N2 H2 DIL. %					
					A	B	A	B	1A	2A	3A	4A	5A	6A	7A	8A	9A	10A	11A	12A			13A	14A	15A		
9-23	3:37	2660	1110	2A	15.31	16.84	13.0	13.2	2.198	2.319	1.907	1.764	1.781	1.764	1.841	1.725	1.818	1.721	1.719	1.819	1.781	1.850	1.972	2.005	2.153	40%	
	4:31	2753	1110	2A	15.29	16.88	13.0	13.1	2.226	2.341	1.909	1.762	1.781	1.758	1.835	1.721	1.804	1.728	1.723	1.824	1.797	1.864	1.995	2.026	2.182	43%	
	5:30	2814	1136	2B	3.34	14.93	4.5	4.5	1.712	1.830	1.696	1.605	1.644	1.615	1.647	1.586	1.637	1.601	1.577	1.676	1.63	1.668	1.688	1.709	1.735	27%	
	6:30	2814	1226	2B	3.44	14.94	4.5	4.5	1.696	1.835	1.705	1.619	1.654	1.628	1.672	1.605	1.658	1.618	1.585	1.679	1.638	1.666	1.671	1.683	1.708	29%	
	8:32	2814	1327	2B	3.46	14.94	4.5	4.5	1.694	1.840	1.708	1.624	1.661	1.634	1.678	1.611	1.663	1.622	1.588	1.681	1.643	1.667	1.671	1.682	1.703	32%	
	8:31	2814	1426	2B	13.48	14.95	4.5	4.5	1.689	1.848	1.709	1.627	1.667	1.638	1.682	1.617	1.670	1.636	1.593	1.683	1.644	1.669	1.673	1.683	1.688	34%	
	9:31	2814	1526	2B	3.51	14.97	4.5	4.5	1.703	1.849	1.707	1.632	1.674	1.647	1.694	1.622	1.678	1.631	1.600	1.687	1.650	1.675	1.679	1.692	1.689	38%	
	10:33	2814	1630	2B	13.57	15.02	4.5	4.5	1.692	1.851	1.709	1.641	1.683	1.655	1.704	1.630	1.691	1.643	1.608	1.693	1.653	1.673	1.669	1.678	1.681	44%	
	11:32	2814	1727	2B	13.58	15.02	4.5	4.5	1.691	1.851	1.708	1.642	1.688	1.659	1.711	1.634	1.694	1.644	1.609	1.695	1.654	1.673	1.668	1.675	1.688	46%	
9-24	12:30	2814	1824	2B	13.59	15.02	4.5	4.5	1.689	1.85	1.708	1.643	1.689	1.661	1.713	1.632	1.693	1.647	1.612	1.697	1.657	1.673	1.667	1.674	1.685	48%	
	1:30	2814	1925	2B	13.61	15.03	4.5	4.5	1.689	1.851	1.708	1.647	1.691	1.663	1.717	1.632	1.692	1.649	1.614	1.697	1.657	1.674	1.669	1.676	1.691	49%	
	2:30	2814	2023	2B	13.61	15.03	4.4	4.5	1.689	1.852	1.707	1.645	1.693	1.664	1.717	1.640	1.702	1.658	1.614	1.700	1.660	1.674	1.668	1.676	1.688	50%	
	3:30	2814	2123	2B	13.58	14.99	4.4	4.4	1.677	1.837	1.704	1.644	1.693	1.665	1.717	1.639	1.702	1.649	1.613	1.688	1.657	1.669	1.661	1.667	1.680	51%	
	4:30	2814	2224	2B	13.62	15.02	4.4	4.4	1.689	1.852	1.712	1.647	1.697	1.668	1.719	1.641	1.703	1.652	1.615	1.701	1.659	1.672	1.663	1.671	1.682	51%	
	5:30			Cal																							
	5:37	2814	2336	2B	13.63	15.05	4.4	4.4	1.689	1.857	1.715	1.647	1.694	1.668	1.718	1.642	1.706	1.657	1.617	1.703	1.661	1.675	1.671	1.676	1.688	51%	
	6:30	2814	2431	2B	13.63	15.05	4.4	4.4	1.689	1.856	1.717	1.645	1.700	1.670	1.718	1.641	1.705	1.655	1.618	1.705	1.661	1.674	1.671	1.677	1.685	51%	
	7:25																										
	7:30																										
	8:35																										
	11:00	2814	2919	2B	15.83	17.32	12.4	12.7	1.977	2.126	2.014	1.866	1.922	1.886	2.007	1.949	1.986	1.937	1.974	1.895	1.911	1.944	1.955	1.988	1.970	71%	
	11:15																										
	12:58	2915	2910	2B	15.10	17.34	13.2	13.3	2.073	2.366	1.976	1.936	1.954	1.822	1.906	1.915	1.912	1.813	1.799	1.874	1.927	1.999	2.038	2.105	2.128	53%	1.92 M
	2:30	2915	2952	2B	15.17	17.40	13.2	13.3	2.102	2.384	1.964	1.932	1.946	1.928	1.913	1.917	1.889	1.863	1.789	1.932	1.868	1.927	2.014	2.054	2.128		
	3:38	3048	3002	2B	13.13	14.67	4.5	4.5	1.738	1.848	1.813	1.594	1.603	1.514	1.604	1.558	1.571	1.564	1.412	1.656	1.64	1.659	1.697	1.732	1.763		
	4:35	3120	3015	2B	15.45	17.10	12.0	12.9	2.139	2.375	1.952	1.810	1.834	1.805	1.881	1.772	1.866	1.784	1.780	1.921	1.854	1.913	2.005	2.060	2.157	54%	
	5:15																										
	5:30	3120	3048	2B	15.39	17.69	13.5	13.5	2.240	2.422	2.000	1.847	1.871	1.836	1.925	1.804	1.904	1.807	1.806	1.949	1.883	1.947	2.057	2.134	2.238	54%	
	6:31	3056	3060	2B	13.12	14.75	4.4	4.4	1.764	1.863	1.693	1.574	1.607	1.575	1.615	1.561	1.586	1.568	1.484	1.656	1.617	1.665	2.021	2.082	2.135	40%	
	6:35																										

\* EVERY 4 HOURS

DATE	TIME	HIGH TIME	LOW TIME	MODE	MOD VOLTS		MODE AMPS		CELL VOLTAGE										FUEL %	MILES																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																	
					A	B	A	B	1A	2A	3A	4A	5A	6A	7A	8A	9A	10A			11A	12A	13A	14A	15A	16A	17A	18A	19A	20A																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																							
9-24	7:31	33:10	30:77	LS15	15.48	14.8																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																															

TEST NO. 1

MODULE STATUS

PAGE 4 OF

DATE	TIME	HIGH TIMER	LOW TIMER	MODE	MOD VOLTS		MOD AMPS		CELL VOLTAGE												EFFICIENT POL	NLMH CONC. *					
					A	B	A	B	1A	2A	3A	4A	5A	6A	7A	8A	9A	10A	11A	12A							
9-25	18:30	4784	57.03	17.15	16.64	18.60	13.3	13.3	2.795	2.515	2.053	1.862	2.132	1.889	1.964	1.840	2.082	1.822	1.838	1.996	1.934	2.002	2.238	2.380	2.453	57.0%	
9-26	17:30	4783	37.15	17.15	18.31	20.5	35.5	35.5	3.18	2.907	2.200	1.915	2.155	2.073	2.111	1.995	2.141	1.858	1.844	2.119	2.035	2.115	2.571	3.017	3.112		
									3.420	6.4	U.S. 411	Ch. 64					back	to	Lo	Mode							
	1:00	4780	38.56	17.1	13.68	15.04	10.76	10.76	1.700	1.725	1.706	1.699	1.966	1.732	1.709	1.624	1.716	1.619	1.593	1.696	1.665	1.647	1.669	1.631	1.697	102.147	342
31	1:30	4780	39.09	17.15	13.73	15.10	10.74	10.74	1.697	1.726	1.713	1.634	1.945	1.732	1.701	1.624	1.727	1.620	1.614	1.705	1.670	1.691	1.672	1.637	1.651	122.0	146
	2:00	4780	37.40	17.15	13.75	15.12	10.71	10.71	1.695	1.724	1.709	1.640	1.921	1.731	1.707	1.623	1.733	1.637	1.618	1.705	1.672	1.691	1.678	1.635	1.705	122.50	146
34	2:30	4780	40.05	17.15	13.82	15.20	10.71	10.71	1.703	1.732	1.721	1.648	1.934	1.732	1.715	1.623	1.736	1.645	1.625	1.714	1.675	1.691	1.672	1.635	1.703	122.86	146
31	3:30	4773	40.27	17.15	13.84	15.19	10.76	10.76	1.694	1.721	1.718	1.636	1.970	1.736	1.725	1.640	1.764	1.652	1.647	1.712	1.678	1.693	1.671	1.631	1.695	12.5	152
31	4:30	4783	42.02	17.15	13.94	15.21	10.76	10.76	1.695	1.723	1.719	1.638	1.933	1.715	1.735	1.644	1.778	1.652	1.627	1.711	1.678	1.691	1.670	1.631	1.692	12.4	206
31	5:30	4783	43.02	17.15	13.95	15.18	10.63	10.63	1.697	1.718	1.714	1.638	2.001	1.712	1.730	1.643	1.774	1.652	1.624	1.710	1.675	1.691	1.668	1.632	1.694	12.6	156
	6:30	4783	43.77	17.15	13.97	15.18	10.51	10.51	1.698	1.722	1.716	1.636	2.008	1.827	1.731	1.641	1.785	1.649	1.625	1.707	1.673	1.688	1.623	1.646	1.697	12.5	156
	7:30																										





DATE		TIME	ELAPSED TIME (HRS)	HYDROGEN OUTPUT METER		POWER INPUT		PRESSURE (PSIG)			COOLANT TEMP (°F)	FLOW RATE		GAS ANALYSIS		METHANOL TEST					
				HIGH VOLTS	LOW VOLTS	HIGH AMPS	LOW AMPS	MOD	P.T.	P.O.	CAL	N <sub>2</sub> H <sub>4</sub>	KCH	QUANT N <sub>2</sub> H <sub>4</sub>	META N <sub>2</sub>		N <sub>2</sub> O <sub>2</sub>	CAPIN	Hz		
10/27	12:43					28.5	5.5	35.5	4.5	157	2.0	5.75	47	3.5	1.7	5.8	3.5	12.3	240	2090	2108
	2:35			28.5	20.5			28.5	4.4	160	2.0	5.60	45	3.3	1.7	5.0	3.5	12.6	255		2117
	4:30					29.5	5.5	26.5	4.4	160	2.0	5.8	45	3.0	1.7	5.5	3.5	12.5	285	2160 cc/min	2126
	6:20	55A																			2136
	8:55																				2143
	9:30	66h				29.5	20.0														
	9:00	66h				29.5	20.0														
	10:00					28.5	5.5	27.5	4.4	156	2.0	5.8	45	4.0	1.7	7.7	3.5	12.5	430	2160 cc/min	2152
	10:30																				
	11:00					N <sub>2</sub> H <sub>4</sub> feed at															
	11:10					N <sub>2</sub> H <sub>4</sub> feed at															
	11:20					N <sub>2</sub> H <sub>4</sub> feed at															
	12:15					BS. 457 cc stop															
	12:30					1047	28.6	20.2	28.7	5.2	27.5	4.3	151	2.0	5.75	46	4.0	1.9	113	2140 cc/min	2162
	2:30					1257	28.6	20.1													
	4:30					1455															
	6:30					1662															
	8:30					1797	28.5	21.2													
	10:30					2011															
	11:00																				
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TEST NO. 2

MODULE STATUS

HRLY

PAGE 1 OF --

DATE	TIME	HIGH TIMER	LOW TIMER	MODE	MOD VOLTS		MOD AMPS		CELL VOLTAGE										EFFLUENT POL	NH <sub>4</sub> CONC. *							
					A	B	A	B	1A	2A	3A	4A	5A	6A	7A	8A	9A	10A			11A	12A					
10/27	12:30	51:26	47:03	H <sub>1</sub>	14.98	16.48	12.7	12.8	1926	1858	1896	1921	1993	1804	1852	1768	1814	1792	1845	1816	1828	1841	1866	1891	590 mm	0.66 (1:27 AM)	
10/27	1:30	52:18	47:39	H <sub>1</sub>	15.13	16.65	12.7	12.8	1923	1871	1916	1932	2.00	1822	1878	1858	1612	1621	1634	1615	1627	1635	1650	1686	460 mm	1.25 (2:34 AM)	
10/27	2:30	52:39	47:58	H <sub>1</sub>	15.27	16.90	12.7	12.8	2008	1917	1919	1942	2.02	1823	1844	1857	1607	1615	1643	1626	1646	1659	1682	1716	380 mm	1.71 (3:30 AM)	
10/27	3:30	52:59	48:16	H <sub>1</sub>	15.34	17.08	12.7	12.8	1711	1673	1654	1637	1670	1601	1626	1573	1603	1616	1647	1627	1645	1660	1679	1709	300 mm	1.89 (4:33 AM)	
"	4:30	53:09	48:40	H <sub>1</sub>	15.50	17.21	12.7	12.9	1996	1943	1937	1964	2.10	1837	1907	1849	1861	1602	1673	1645	1656	1670	1691	1722	380 mm	1.07 (5:45 AM)	
"	5:30	53:21	48:57	H <sub>1</sub>	15.54	17.26	12.7	12.9	2009	1951	1938	1975	2.10	1840	1911	1848	1870	1831	1835	1901	1918	1944	1975	2002	510 mm	1.07 (6:45 AM)	
"	6:30	54:32	49:54	H <sub>1</sub>	15.60	17.41	12.7	12.9	2010	1959	1946	1983	2.12	1841	1917	1854	1887	1837	1937	1902	1915	1934	1958	1992	510	510	1.07 (7:45 AM)
"	7:30	54:47	49:25	H <sub>1</sub>	15.64	17.49	12.7	12.9	2010	1959	1946	1983	2.12	1841	1917	1854	1887	1837	1937	1902	1915	1934	1958	1992	510	510	1.07 (8:45 AM)
"	8:30	55:54	49:59	H <sub>1</sub>	15.60	17.29	12.8	12.9	2000	1951	1946	2008	2.14	1850	1932	1805	1867	1804	1841	1857	1915	1932	1965	2001	2012	510 mm	1.07 (9:45 AM)
"	9:30	55:54	49:59	H <sub>1</sub>	15.64	17.49	12.8	13.0	2058	1982	1947	2020	2.14	1848	1902	1863	1887	1846	1870	1862	1870	1886	1902	1917	510	510	1.07 (10:45 AM)
10-28	12:00	N/A	N/A	"	17.4	much less																					
10-28	12:30	57:46	50:68	H <sub>1</sub>	15.71	17.57	12.8	13.0	2058	1982	1947	2020	2.14	1848	1902	1863	1887	1846	1870	1862	1870	1886	1902	1917	510	510	1.07 (11:45 AM)
"	1:30	58:03	50:31	H <sub>1</sub>	15.74	17.63	12.8	13.0	2078	1985	1948	2020	2.14	1848	1902	1863	1887	1846	1870	1862	1870	1886	1902	1917	510	510	1.07 (12:45 AM)
"	2:30	58:25	51:14	H <sub>1</sub>	15.76	17.67	12.8	13.0	2085	1999	1948	2027	2.14	1848	1902	1863	1887	1846	1870	1862	1870	1886	1902	1917	510	510	1.07 (1:45 AM)
"	3:30	59:08	51:57	H <sub>1</sub>	15.71	17.76	12.7	12.9	2011	2011	1947	2011	2.14	1848	1902	1863	1887	1846	1870	1862	1870	1886	1902	1917	510	510	1.07 (2:45 AM)
"	4:30	59:20	51:57	H <sub>1</sub>	15.71	17.68	12.7	12.9	2011	2011	1947	2011	2.14	1848	1902	1863	1887	1846	1870	1862	1870	1886	1902	1917	510	510	1.07 (3:45 AM)
"	5:30	60:17	51:57	H <sub>1</sub>	15.71	17.68	12.7	12.9	2011	2011	1947	2011	2.14	1848	1902	1863	1887	1846	1870	1862	1870	1886	1902	1917	510	510	1.07 (4:45 AM)
"	6:30	60:38	52:17	H <sub>1</sub>	15.58	17.61	12.8	13.0	2042	1980	1953	2020	2.10	1846	1894	1873	1846	1854	1880	1854	1867	1886	1902	1917	510	510	1.07 (5:45 AM)
"	7:30	61:42	52:54	H <sub>1</sub>	15.68	17.87	12.8	13.0	2049	2007	1949	2022	2.09	1835	1875	1783	1835	1854	1854	1854	1854	1854	1854	1854	510	510	1.07 (6:45 AM)
"	8:30	61:58	52:54	H <sub>1</sub>	15.67	17.76	12.8	13.0	2049	2007	1949	2022	2.09	1835	1875	1783	1835	1854	1854	1854	1854	1854	1854	1854	510	510	1.07 (7:45 AM)
"	9:30	62:16	53:24	H <sub>1</sub>	15.63	17.83	12.8	13.0	2049	2007	1949	2022	2.09	1835	1875	1783	1835	1854	1854	1854	1854	1854	1854	1854	510	510	1.07 (8:45 AM)
"	10:30	62:16	53:24	H <sub>1</sub>	15.48	17.87	12.8	13.0	2049	2007	1949	2022	2.09	1835	1875	1783	1835	1854	1854	1854	1854	1854	1854	1854	510	510	1.07 (9:45 AM)
"	11:30	63:16	53:24	H <sub>1</sub>	15.48	17.87	12.8	13.0	2049	2007	1949	2022	2.09	1835	1875	1783	1835	1854	1854	1854	1854	1854	1854	1854	510	510	1.07 (10:45 AM)
"	12:30	63:16	53:24	H <sub>1</sub>	15.48	17.87	12.8	13.0	2049	2007	1949	2022	2.09	1835	1875	1783	1835	1854	1854	1854	1854	1854	1854	1854	510	510	1.07 (11:45 AM)
"	1:30	64:15	54:30	H <sub>1</sub>	15.56	18.11	12.8	13.0	2049	2007	1949	2022	2.09	1835	1875	1783	1835	1854	1854	1854	1854	1854	1854	1854	510	510	1.07 (12:45 AM)
"	2:30	64:15	54:30	H <sub>1</sub>	15.56	18.11	12.8	13.0	2049	2007	1949	2022	2.09	1835	1875	1783	1835	1854	1854	1854	1854	1854	1854	1854	510	510	1.07 (1:45 AM)
"	3:30	65:01	55:08	H <sub>1</sub>	15.41	18.09	12.8	13.0	2049	2007	1949	2022	2.09	1835	1875	1783	1835	1854	1854	1854	1854	1854	1854	1854	510	510	1.07 (2:45 AM)
4:10				N <sub>2</sub> makeup																							
4:25				H <sub>2</sub> O																							

\* EVERY 4 HOURS

TEST NO. 2

MODULE STATUS

HRLY

PAGE 2 OF

DATE	TIME	HIGH TIMER	LOW TIMER	MODE	MOD VOLTS		MOD AMPS		CELL VOLTAGE										EFFLUENT POL	NZHA CONC. *
					A	B	A	B	1A	2A	3A	4A	5A	6A	7A	8A	9A	10A	11A	12A
10/28	4:30	6564	5539	H <sub>1</sub>	15.28	18.28	12.8	13.1	2.117	1.561	1.951	2.006	1.777	1.771	1.731	1.782	1.843	1.805	1.952	1.948
"	5:30	6610	5576	L <sub>1</sub>	13.91	15.90	4.3	4.3	1.750	1.697	1.673	1.692	1.702	1.604	1.615	1.578	1.611	1.641	1.609	1.681
"	6:30	6651	5594	H <sub>1</sub>	15.83	17.79	12.8	13.0	2.074	2.021	1.983	2.078	2.140	1.864	1.811	1.872	1.898	1.862	2.013	1.973
"	7:30	6712	5634	H <sub>1</sub>	15.85	18.04	4.3	4.3	1.761	1.731	1.678	1.689	1.739	1.627	1.643	1.593	1.632	1.652	1.708	1.680
"	8:30	6759	5680	L <sub>1</sub>	13.30	15.31	4.3	4.3	1.740	1.717	1.672	1.681	1.706	1.620	1.631	1.589	1.627	1.649	1.708	1.680
"	9:30	6806	5702	L <sub>1</sub>	13.33	15.23	4.3	4.3	1.737	1.709	1.674	1.675	1.705	1.620	1.633	1.583	1.621	1.643	1.708	1.680
"	10:30	6859	5744	H <sub>1</sub>	15.78	17.97	12.8	13.0	2.067	2.001	1.987	2.074	2.118	1.857	1.893	1.808	1.871	1.860	2.017	1.979
"	11:30	6912	5776	H <sub>1</sub>	15.69	17.96	12.9	13.1	2.075	2.009	1.991	2.074	2.121	1.858	1.893	1.808	1.868	1.863	2.018	1.982
10/29	12:30	6960	5812	L <sub>1</sub>	13.33	15.24	4.35	4.4	1.727	1.727	1.663	1.644	1.648	1.601	1.605	1.553	1.605	1.587	1.717	1.630
"	"	"	"	H <sub>1</sub>	15.36	18.37	"	"	2.165	2.094	1.973	2.060	2.094	1.893	"	"	1.857	1.803	2.049	1.984
"	1:30	7004	5852	H <sub>1</sub>	15.31	18.01	12.9	13.2	2.082	2.014	1.987	2.073	2.115	1.858	1.890	1.804	1.857	1.803	2.049	1.984
"	2:30	7033	5883	H <sub>1</sub>	15.31	18.48	12.9	13.2	2.169	2.080	1.980	2.083	2.083	"	"	"	1.857	1.803	2.049	1.984
"	"	"	"	"	13.16	15.33	4.3	4.4	1.771	1.717	1.668	1.653	1.639	1.584	1.574	1.545	1.600	1.631	1.580	1.717
"	3:30	7150	5906	H <sub>1</sub>	15.81	18.39	12.9	13.2	2.161	2.097	1.982	2.083	2.084	1.849	1.875	1.799	1.852	1.804	2.049	1.984
"	4:30	7222	5939	H <sub>1</sub>	15.74	18.12	12.9	13.2	2.079	2.006	1.980	2.083	2.083	1.849	1.875	1.799	1.852	1.804	2.049	1.984
"	5:30	7267	6038	H <sub>1</sub>	15.75	18.28	12.9	13.2	2.102	2.035	1.996	2.083	2.091	1.839	1.862	1.795	1.840	1.801	1.872	2.049
"	6:30	7338	6061	L <sub>1</sub>	13.01	15.33	4.4	4.3	1.772	1.709	1.667	1.654	1.639	1.574	1.574	1.533	1.580	1.628	1.601	1.715
"	7:30	7373	6106	H <sub>1</sub>	13.32	15.42	12.8	13.2	2.090	2.016	1.973	1.669	1.654	1.580	1.589	1.552	1.589	1.622	1.899	1.698
"	8:30	7435	6138	H <sub>1</sub>	13.37	15.33	12.8	13.2	1.960	1.907	1.685	1.699	1.720	1.616	1.622	1.576	1.616	1.640	1.611	1.701
"	9:30	7499	6178	H <sub>1</sub>	15.93	18.19	12.8	13.2	2.057	2.004	2.011	2.133	2.164	1.854	1.895	1.808	1.874	1.897	1.861	2.022
"	10:30	7514	6216	H <sub>1</sub>	15.98	18.20	12.9	13.2	2.059	2.009	2.027	2.142	2.172	1.869	1.912	1.808	1.887	1.908	1.873	2.037
"	11:30	7564	6259	H <sub>1</sub>	15.84	18.21	12.8	13.2	2.079	1.981	2.019	2.125	2.155	1.843	1.881	1.784	1.856	1.888	1.850	1.978
"	12:30	7623	6287	H <sub>1</sub>	15.90	18.41	12.8	13.2	2.109	2.012	2.033	2.133	2.146	1.847	1.897	1.807	1.864	1.890	1.858	2.027
"	1:30	7686	6308	H <sub>1</sub>	15.76	18.48	12.8	13.1	2.124	1.994	2.009	2.133	2.146	1.847	1.897	1.807	1.864	1.890	1.858	2.027
"	2:30	7728	6344	H <sub>1</sub>	15.75	18.28	12.8	13.0	2.105	1.998	2.019	2.093	2.116	1.820	1.846	1.747	1.817	1.833	1.811	1.928
"	3:30	7760	6379	H <sub>1</sub>	15.69	18.01	12.8	13.1	2.089	1.994	2.012	2.094	2.115	1.827	1.851	1.748	1.833	1.872	1.838	1.992
"	4:30	7760	6379	H <sub>1</sub>	15.69	18.01	12.8	13.1	2.089	1.994	2.012	2.094	2.115	1.827	1.851	1.748	1.833	1.872	1.838	1.992

\* EVERY 4 HRS

TEST NO. 2

MODULE STATUS

PAGE 3 OF

3252 3309

DATE	TIME	HIGH TIMER	LOW TIMER	MODE	MOD VOLTS		MOD AMPS		CELL VOLTAGE										EFFECTIVE POL	NEMA CORRECTION *								
					A	B	A	B	1A	2A	3A	4A	5A	6A	7A	8A	9A	8B			7B	6B	5B	4B	3B	2B	1B	
10/29	5:30	78.18	64.15	64.15	15.58	18.26	12.8	13.0	2.102	1.982	1.998	2.059	2.082	1.798	1.815	1.732	1.607	1.429	1.802	1.935	1.915	2.057	2.181	2.381	2.432	460		
	6:55	78.96	64.49	64.49																								
	7:25	79.17	64.69	64.69	15.54	18.59	13.0	13.4	2.133	2.030	2.002	2.006	2.097	1.843	1.864	1.787	1.844	1.897	1.868	2.062	2.017	2.077	2.211	2.364	2.379	460		
	8:30	79.88	64.99	64.99	15.92	18.91	12.9	13.1	2.142	2.034	2.002	2.106	2.117	1.856	1.877	1.993	1.865	1.898	1.867	2.054	1.990	2.050	2.182	2.318	2.354	465		
	9:30	80.52	65.32	65.32	13.25	15.28	4.3	4.4	1.765	1.714	1.674	1.657	1.641	1.610	1.611	1.587	1.607	1.692	1.610	1.570	2.035	2.009	2.055	2.113	2.176	450		
	10:30	81.07	65.62	65.62	15.59	18.32	13.0	13.3	2.149	2.020	2.005	2.066	2.031	1.837	1.848	1.767	1.826	1.881	1.955	2.046	2.032	2.102	2.188	2.339	2.356	430		
10/30	12:00			N <sub>2</sub> H <sub>4</sub>																								
	12:30	82.12	66.20	66.20	15.82	19.06	13.1	13.3	2.145	2.043	2.015	2.034	2.033	1.833	1.852	1.776	1.812	1.892	1.855	2.035	2.046	2.135	2.231	2.463	2.519	420		
	1:30	82.63	66.46	66.46	15.91	19.75	13.0	13.1	2.164	2.040	2.018	2.003	2.000	1.856	1.869	1.798	1.831	1.893	1.863	2.000	2.030	2.105	2.265	2.407	2.450	450		
	2:30	83.31	66.76	66.76	15.84	19.78	13.0	13.4	2.145	2.025	2.007	2.009	1.943	1.853	1.786	1.824	1.891	1.859	2.001	2.041	2.129	2.230	2.448	2.520	450			
	3:00	83.84	67.08	67.08	13.13	15.32	4.3	4.4	1.770	1.707	1.672	1.657	1.647	1.577	1.594	1.554	1.586	1.635	1.595	1.706	1.676	1.746	1.757	1.792	1.817	255		
	4:30	84.25	67.39	67.39	16.0	19.48	13.0	13.3																				
	5:30	84.85	67.75	67.75	15.98	19.28	13.2	13.4	2.210	2.051	2.045	2.097	2.017	1.851	1.862	1.799	1.835	1.900	1.867	2.111	2.062	2.156	2.360	2.498	2.580	430		
	6:30	85.35	68.08	68.08	15.92	18.93	13.1	13.3	2.192	2.034	2.034	2.101	2.001	1.848	1.862	1.780	1.823	1.891	1.858	2.003	2.038	2.113	2.301	2.447	2.505	430		
	6:58	N <sub>2</sub> H <sub>4</sub>	Flow 0	11.0									6.955															
	7:16	O <sub>2</sub> 5.65																										
	7:30	85.91	68.30	68.30	15.92	19.32	13.2	13.4	2.222	2.049	2.029	2.101	2.060	1.829	1.849	1.782	1.807	1.877	1.851	2.079	2.059	2.187	2.420	2.633	2.678	260		
	8:30	86.59	68.65	68.65	15.88	18.79	13.0	13.4	2.191	2.029	2.028	2.080	2.072	1.838	1.858	1.786	1.821	1.896	1.857	2.094	2.057	2.145	2.320	2.493	2.541	420		
	9:30	86.97	68.98	68.98	15.59	18.24	13.0	13.2	2.172	2.033	2.032	2.103	2.079	1.839	1.855	1.788	1.817	1.887	1.854	2.083	2.052	2.143	2.320	2.494	2.547	430		
	10:30	87.53	69.12	69.12	12.57	19.04	4.5	4.5	1.784	1.705	1.661	1.587	1.576	1.905	1.834	1.807	1.531	1.850	1.820	1.998	2.007	2.107	2.181	2.276	2.304	410		

avg - 2.055

\* EVERY 4 HOURS



TEST NO. 3

SYSTEM STATUS

2 hr

PAGE 1 OF —

DATE	TIME	ELAPSED TIME (HRS)	HYDROGEN OUTPUT METER	POWER INPUT		PRESSURE (PSIG)			COOLANT TEMP (°F)	FLOW RATE			GAS ANALYSIS					
				HIGH VOLTS	LOW VOLTS	MOD	PI	PD2		CAL	NH4	KOH	COALANT IN-H4	META	N2	BUB °F	N2-O2	CABIN
11-17	10:08		N2H4	feed on	at 20.5		10.08	2.0	1.9	5.9	high @ 11.055	500 on valve						
	10:30		221			41	4.5	152	1.9	5.9	5.5	11.055	4.25	16.55	45	2720 cc/min		222x1
am	11:30		218			28.5	5.5	43.5	4.5	151	1.9	5.9	5.5	16.55	45	2720 cc/min		2227
pm	12:30	0																2233
	12:45	45	430			28.5	5.5	46.5	4.5	145	1.9	5.8	6.2	12.5	50	2130 cc/min		
	1:06	1:06	N2H4	feed on	at 20.5		10.08	2.0	1.9	5.9	high @ 11.055	500 on valve						
	1:55	1:55	550			34.5	4.5	141	1.9	5.8	47	7.8	7.7	12.6	70	2130 cc/min		
	4:05	4:05	774			38.5	4.3	142	1.9	5.9	47	6.3	6.3	12.5	85			
	4:45	4:45				3.35	min	2.2	10	2.0	46	2190						
	5:58					N2H4 on		Value blank	open	6:10 with flow								
	6:17					10	cc	N2H4 sample										78M
	6:23	6:23	980			28.6	20.0	27	4.3	155	1.9	5.8	47	12.5	105			
	8:00	8:00	1108			28.6	5.5	31	4.3	157	1.9	5.8	46	12.5	110	2190 cc/min		2251
	8:30	Note	gain			2 in	in	no feed	in 1 hr									
	8:30	Note	N2H4 in 30 min			1 in	in	Reduce flow	to 390	to 1 hr								
	7:30	Note	N2H4 in 30 min			1 in	in	Reduce flow	to 390	to 1 hr								
	10:00	10:00	1348			28.7	20.8	28.4	5.2	3.5	4.3	154	1.9	5.8	43			2260
	10:10	Note	low			new	new	new	new	new	new	new	new	new	new			
	11:30	Note	low			new	new	new	new	new	new	new	new	new	new			
mid	12:00	12	1591			20.5	5.5	32	4.5	150	1.9	5.8	42	12.5	163			
am	2:00	14	1625			28.5	20.0	37.5	4.5	147	1.9	5.8	42	12.5	187			
	3:16					Reduce	N2H4 flow	to 390	to 390	to 390	to 390	to 390	to 390	to 390	to 390			
	3:25		down N2H4 tank			add	600 cc	to 600 cc	to 600 cc	to 600 cc	to 600 cc	to 600 cc	to 600 cc	to 600 cc	to 600 cc			
	4:02	16	1617			28.5	5.8	27	4.5	144	1.9	5.8	42	12.5	225			
	4:30					8.5	at 28.0 cc											
	6:00	18	2092			20.5	20.5	36	4.5	152	1.9	5.8	42	12.5	550			
	8:00	20	2391			28.5	5.5	34	4.5	160	1.8	5.6	43	12.4	85			
	8:50	21	Bubble			separator	level raised	from 100	to 75									
	10:00	22	2589			20.5	21.0	31	4.5	157	1.6	5.75	42	12.4	95			
	10:15		Bubble			separator	level raised	from 100	to 75									





TEST NO. 3

MODULE STATUS

PAGE 1 OF

DATE	TIME	HIGH TIMER	LOW TIMER	MODE	MOD VOLTS		MOD AMPS		CELL VOLTAGE																EFFECTIVE POL	NLMA COST		
					A	B	A	B	1A	2A	3A	4A	5A	6A	7A	8A	9A	8B	7B	6B	5B	4B	3B	2B			1B	
11-10	10:36	9362	6960	H1	15.18	17.01	12.75	12.90	1.995	1.916	1.897	1.877	1.931	1.830	1.928	1.824	1.830	1.935	1.815	1.914	1.857	1.995	1.934	1.948	1.994	640	10.98(00)	
	11:35	9426	6950	H1	15.19	17.24	12.80	13.0	2.065	1.954	1.901	1.887	1.905	1.815	1.893	1.807	1.810	1.943	1.821	1.951	1.891	1.938	1.985	2.008	2.073	530	11.04(00)	
	2:05	9510	7047	L0	13.09	14.85	4.3	4.3	1.695	1.676	1.666	1.625	1.605	1.624	1.666	1.616	1.628	1.861	1.833	1.867	1.915	1.948	1.980	1.991	2.037	440	12.04(00)	
	3:00	9532	7076	L0	13.18	14.90	4.3	4.3	1.634	1.629	1.621	1.642	1.623	1.604	1.638	1.582	1.601	1.648	1.537	1.612	1.505	1.532	1.567	1.591	2.042	350		
	4:00	9586	7111	H1	15.52	17.58	13.0	13.2	2.067	1.994	1.954	1.941	2.008	1.848	1.912	1.818	1.857	1.874	1.833	1.939	1.969	2.012	2.045	2.084	2.140	540		
	5:00	9615	7127	H1	15.63	17.75	13.0	13.2	2.081	2.016	1.961	1.961	2.031	1.848	1.907	1.823	1.865	1.801	1.854	1.907	1.946	1.974	2.067	2.112	2.162	570		
	6:00	9616	7132	H1																								
	6:20	9634	7154	H1	15.64	17.85	13.0	13.2	2.083	2.016	1.962	1.962	2.033	1.841	1.892	1.823	1.865	1.804	1.851	1.907	1.932	1.935	2.027	2.094	2.136	590	7.84(17-100)	
	7:00	9736	7173	H1	15.55	17.86	13.0	13.2	2.098	2.033	1.963	1.947	1.975	1.831	1.871	1.816	1.857	1.833	1.857	2.016	1.938	1.952	2.033	2.110	2.149	560		
	8:00	9115	7200	H1	15.5	18.07	13.1	13.3	2.132	2.066	1.957	1.937	1.972	1.812	1.842	1.806	1.856	1.830	1.853	1.711	1.607	1.705	1.768	1.771	1.721	510		
	9:00	9846	7236	H1	15.48	17.84	13.1	13.3	2.014	2.1	1.966	1.952	1.948	1.844	1.864	1.811	1.826	1.821	1.848	2.020	1.770	2.002	2.040	2.125	2.146	480		
	10:00	9918	7254	H1	15.53	18.43	13.2	13.3	2.225	2.117	1.942	1.944	1.944	1.785	1.830	1.776	1.785	1.870	1.847	2.451	2.008	2.410	2.177	2.320	2.412	470		
	11:00	9935	7281	L0	12.17	15.03	4.6	4.5	1.768	1.729	1.666	1.638	1.605	1.400	1.511	1.70	1.541	1.540	1.449	2.438	1.944	2.438	2.018	2.143	2.114	160/48		
11-11	12:00	9985	7300	H1	14.61	17.10	13.2	13.4	2.256	2.177	1.980	1.913	1.903	1.793	1.837	1.799	1.819	1.885	1.855	2.103	2.045	2.107	2.195	2.286	2.481	410		
	1:00	10035	7320	L0	12.34	14.97	4.6	4.5	1.816	1.733	1.678	1.630	1.608	1.219	1.506	1.397	1.538	1.591	1.544	1.699	1.678	1.775	1.743	1.792	1.819	170		
	2:00	10143	7356	H1	15.66	18.22	12.9	13.2	2.211	2.133	1.976	1.967	1.967	1.801	1.851	1.805	1.871	1.841	2.066	2.013	2.106	2.156	2.237	2.408	15			
	3:00	10140	7383	H1	15.77	18.80	13.1	13.4	2.284	2.207	1.972	1.947	1.922	1.803	1.899	1.785	1.787	1.843	1.851	2.110	2.054	2.115	2.225	2.449	2.555	120		
	4:00	10213	7410	H1	15.98	19.11	13.2	13.4	2.300	2.233	1.969	1.934	1.919	1.619	1.672	1.547	1.508	1.545	1.578	1.718	1.691	1.744	1.767	1.817	1.849	2.15		
	5:00	10266	7443	L0	15.71	18.03	12.9	13.2	2.155	2.081	1.986	1.994	2.083	1.863	1.878	1.805	1.812	1.873	1.844	2.040	2.001	2.053	2.131	2.223	2.362	530		
	6:00	10321	7462	H1	15.75	18.50	12.9	13.1	2.195	2.069	1.987	1.974	2.063	1.844	1.869	1.779	1.807	1.849	1.861	1.833	1.933	2.044	2.186	2.387	2.512			
	7:00	10384	7475	H1	15.94	18.98	13.1	13.4	2.233	2.120	2.000	2.017	2.078	1.841	1.866	1.805	1.829	1.861	1.833	2.061	2.011	2.127	2.243	2.483	2.550	550		
	8:00	10455	7504	H1	15.68	18.62	13.1	13.6	2.195	2.085	1.991	1.984	2.004	1.806	1.831	1.789	1.812	1.866	1.847	2.095	2.048	2.156	2.372	2.584	2.591	420		
	9:00	10517	7528	H1	15.72	18.63	13.1	13.5	2.242	2.116	1.984	1.976	1.995	1.809	1.825	1.781	1.802	1.854	1.837	2.072	2.020	2.134	2.308	2.437	2.445	460		
	10:00	10589	7563	H1	15.70	18.41	13.1	13.5	2.244	2.128	2.011	1.964	1.952	1.804	1.829	1.784	1.804	1.862	1.837	2.028	1.995	2.095	2.176	2.311	2.401	450		
	11:00	10655	7582	H1	15.62	18.80	13.1	13.5	2.267	2.089	1.975	1.922	1.977	1.820	1.791	1.750	1.777	1.825	1.805	1.993	1.967	2.142	2.342	2.505	2.602	420		
	12:00	10730	7613	L0	12.22	17.97	4.0	3.8	2.232	2.067	1.980	1.945	1.962	1.835	1.813	1.780	1.811	1.843	1.821	2.024	2.003	2.118	2.251	2.341	2.372	450		
	1:00	10794	7630	L0	11.58	14.83	4.0	3.8	1.837	1.755	1.687	1.600	1.454	1.407	1.408	1.391	1.505	1.549	1.491	1.694	1.674	1.732	1.757	1.788	1.805	140		
	2:00	10845	7652	L0	11.57	14.83	4.0	3.8	2.193	2.101	2.007	1.991	2.059	1.835	1.861	1.838	1.808	1.864	1.838	2.049	2.008	2.073	2.172	2.262	2.336	440		
	3:00	10896	7686	L0	13.09	15.15	4.3	4.4	2.181	2.054	1.972	1.966	2.045	1.822	1.855	1.802	1.794	1.845	1.822	2.005	1.981	2.095	2.222	2.338	2.374	490		
	4:00	10945	7716	H1	16.03	18.52	2.9	13.6	2.207	2.103	1.997	1.995	2.096	1.831	1.862	1.829	1.829	1.816	1.821	1.977	1.957	2.055	2.179	2.352	2.508	520		

TEST NO. 3

MODULE STATICS

PAGE 2 JF

DATE	TIME	HIGH TIMER	LOW TIMER	MODE	MOD VOLTS		MOD AMPS		CELL VOLTAGE										VOLTAGE POT.	NEMA CODE						
					A	B	A	B	1A	2A	3A	4A	5A	6A	7A	8A	9A	10A			11A	12A	13A	14A	15A	16A
11-11	5:00	10:02	77:54	H1	16.2	15.39	13.9	13.0	2.157	2.106	2.077	2.026	2.136	2.000	1.913	1.957	1.953	1.871	1.935	2.047	1.971	2.030	2.157	2.278	2.365	550
	6:00	10:26	77:16	H1	16.14	15.34	13.1	13.3	2.145	2.077	2.006	2.030	2.146	1.993	1.917	1.959	1.960	1.881	1.945	2.045	1.978	2.041	2.138	2.230	2.343	580
	7:00	10:42	78:04	H1	16.25	15.45	13.1	13.3	2.200	2.104	2.002	2.036	2.148	1.994	1.916	1.952	1.956	1.870	1.940	2.040	1.973	2.040	2.138	2.230	2.343	570
	8:00	10:56	78:53	H1	16.09	15.52	13.1	13.3	2.225	2.095	1.999	2.030	2.129	1.955	1.880	1.932	1.934	1.867	1.935	2.040	1.973	2.040	2.138	2.230	2.343	570
	9:00	11:03	78:51	H1	16.21	15.52	13.1	13.3	2.225	2.095	1.999	2.030	2.129	1.955	1.880	1.932	1.934	1.867	1.935	2.040	1.973	2.040	2.138	2.230	2.343	570
	10:00	11:14	79:00	H1	16.16	15.51	13.2	13.4	2.233	2.140	2.033	2.045	2.108	1.914	1.870	1.964	1.917	1.804	1.946	2.040	1.973	2.040	2.138	2.230	2.343	570
	11:00	11:27	79:18	H1	16.32	15.40	13.5	13.7	2.306	2.215	2.098	2.109	2.169	1.910	1.864	1.958	1.911	1.800	1.946	2.040	1.973	2.040	2.138	2.230	2.343	570
11-12	12:00	11:34	79:55	H1	16.27	15.40	13.4	13.6	2.392	2.202	2.098	2.109	2.169	1.910	1.864	1.958	1.911	1.800	1.946	2.040	1.973	2.040	2.138	2.230	2.343	570
	1:00	12:03	79:57	LO	11.87	11.43	4.6	4.6	1.855	2.080	1.988	2.073	2.100	1.991	1.856	1.852	1.844	1.846	1.822	1.937	1.962	2.100	2.247	2.361	2.492	420
	2:00	14:53	80:04	LO	13.17	12.79	4.5	4.5	1.843	1.720	1.664	1.621	1.620	1.591	1.527	1.525	1.524	1.580	1.550	1.659	1.653	1.737	1.789	1.820	1.895	220
	3:00	15:04	80:06	H1	16.49	15.24	4.4	4.4	1.859	1.746	1.699	1.663	1.707	1.670	1.613	1.597	1.617	1.621	1.577	1.663	1.652	1.735	1.752	1.810	1.850	380
	4:00	15:10	80:06	H1	16.57	15.27	4.4	4.4	2.348	1.769	1.721	1.682	1.780	1.702	1.639	1.609	1.632	1.618	1.586	1.671	1.654	1.718	1.738	1.805	1.843	480
	5:00	16:34	80:07	H1	16.53	15.27	4.4	4.4	2.271	2.095	2.061	2.072	2.157	2.055	1.917	1.889	1.937	1.859	1.820	1.923	1.923	1.727	1.753	1.818	1.887	570
	6:00	16:55	81:05	H1	16.52	15.27	4.4	4.4	2.223	2.087	2.049	2.079	2.165	2.080	1.938	1.896	1.941	1.873	1.841	1.923	1.923	1.727	1.753	1.818	1.887	610
	7:00	17:51	81:55	LO	13.08	12.45	4.2	4.2	1.813	1.726	1.703	1.695	1.657	1.747	1.657	1.636	1.653	1.632	1.602	1.671	1.651	1.711	1.729	1.803	1.837	560
	8:00	18:01	81:54	H1	16.70	15.01	4.3	4.3	1.849	1.752	1.709	1.689	1.761	1.703	1.645	1.614	1.644	1.619	1.587	1.678	1.667	1.721	1.743	1.809	1.845	420
	9:00	18:08	82:24	H1	16.58	15.76	4.4	4.4	2.273	2.109	2.056	2.113	2.169	2.033	1.924	1.879	1.917	1.877	1.834	1.916	1.916	2.078	2.122	2.193	2.230	520
	10:00	19:05	82:25	H1	16.46	15.71	4.3	4.3	2.231	2.122	2.038	2.084	2.162	2.014	1.915	1.878	1.934	1.880	1.837	1.921	1.921	2.078	2.122	2.193	2.230	520
	11:00	19:38	83:09	H1	16.43	15.80	4.3	4.3	2.326	2.147	2.016	2.048	2.134	1.971	1.906	1.858	1.879	1.864	1.824	1.903	1.903	1.731	1.768	1.831	1.853	400
	12:00	20:05	83:26	LO	12.18	11.49	4.3	4.3	1.890	1.761	1.699	1.512	1.264	1.942	1.471	1.112	1.231	1.434	1.408	1.674	1.658	1.733	1.762	1.833	1.870	150
	1:00	20:34	83:26	LO	11.84	11.43	4.6	4.6	1.873	1.776	1.698	1.591	1.283	1.972	1.492	1.121	1.125	1.408	1.371	1.665	1.654	1.725	1.761	1.833	1.870	130
	2:30	21:15	84:02	LO	12.14	11.49	4.5	4.5	1.864	1.734	1.673	1.621	1.585	1.936	1.494	1.187	1.529	1.558	1.471	1.647	1.643	1.719	1.753	1.827	1.857	170
	3:00	21:49	84:22	H1	15.73	17.26	12.9	13.2	2.208	2.034	1.946	1.948	2.084	1.911	1.847	1.805	1.856	1.839	1.793	1.813	1.808	1.831	1.811	1.844	1.874	470
	4:00	22:25	84:40	H1	16.33	19.14	13.2	13.4	2.390	2.081	1.994	1.988	2.110	2.008	1.887	1.851	1.955	1.853	1.803	1.964	1.921	2.075	2.034	2.116	2.144	535
11-13	5:00	22:58	84:44	H1	16.64	19.17	13.4	13.4	2.405	2.117	2.026	2.022	2.124	2.062	1.935	1.867	1.974	1.849	1.809	1.988	1.913	2.036	2.032	2.116	2.144	570
	6:00	23:11	85:31	LO	13.65	15.37	4.3	4.3	1.830	1.700	1.711	1.617	1.775	1.654	1.632	1.614	1.657	1.639	1.599	1.649	1.624	1.720	1.734	1.796	1.844	380/450
	7:00	23:41	85:31	LO	13.65	15.37	4.3	4.3	1.830	1.700	1.711	1.617	1.775	1.654	1.632	1.614	1.657	1.639	1.599	1.649	1.624	1.720	1.734	1.796	1.844	380/450
	8:00	24:11	85:31	H1	16.53	19.07	13.1	13.2	2.371	2.104	2.032	2.037	2.152	2.047	1.934	1.878	1.983	1.854	1.835	2.092	2.032	2.141	2.149	2.239	2.267	520
	9:00	24:51	85:33	H1	16.64	19.42	13.2	13.3	2.396	2.181	2.031	2.037	2.142	2.027	1.933	1.887	1.984	1.854	1.835	2.092	2.032	2.141	2.149	2.239	2.267	520
	10:00	25:48	85:30	LO	12.34	14.77	4.3	4.3	1.852	1.757	1.653	1.653	1.593	1.331	1.425	1.448	1.528	1.549	1.454	1.615	1.623	1.734	1.750	1.810	1.844	460
	11:00	26:55	86:30	H1	16.71	19.15	13.3	13.3	2.370	2.177	2.027	2.037	2.142	2.027	1.933	1.887	1.984	1.854	1.835	2.092	2.032	2.141	2.149	2.239	2.267	520
	12:00	27:14	86:58	LO	11.74	14.69	4.6	4.6	2.347	2.131	2.042	2.111	2.021	1.928	1.858	1.800	1.900	2.003	1.881	2.005	1.998	2.117	2.117	2.200	2.200	430
	1:00	27:44	86:58	H1	16.74	19.34	13.4	13.4	2.378	2.161	2.044	2.044	2.142	2.027	1.933	1.887	1.984	1.854	1.835	2.092	2.032	2.141	2.149	2.239	2.267	520

\* EVERY 4 HOURS



DATE	TIME	HIGH TIMER	LOW TIMER	MODE	MOD VOLTS		MOD AMPS		CELL VOLTAGE												EFFICIENT POL	NLMA CORIC. *					
					A	B	A	B	1A	2A	3A	4A	5A	6A	7A	8A	9B	8B	7B	6B			5B	4B	3B	2B	1B
11-13	2:00	2723	8704	LO	15.81	15.50	4.3	4.4	1837	1750	1717	1668	1623	1607	1624	1601	1639	1611	1576	1665	1647	1715	1730	1791	1842	350	
	3:00	2762	8737	HI	16.64	15.11	4.3	4.4	1832	1762	1707	1691	1604	1757	1698	1624	1662	1622	1584	1659	1638	1684	1699	1786	1838	470	
	4:00	2815	8757	HI	16.76	15.03	4.3	4.4	2363	2121	2081	2102	2150	2094	1966	1909	2021	1884	1831	2022	1939	2075	2342	2476	2610	590	83M
	5:00	2859	8784	HI	16.95	15.30	4.3	4.4	2360	2104	2077	2102	2150	2094	1966	1909	2021	1884	1831	2022	1939	2075	2342	2476	2610	470	
	6:00	2907	8810	HI	16.42	18.93	4.3	4.4	2332	2085	2073	2052	2114	2034	1927	1920	2011	1889	1806	1986	1938	2103	2364	2524	2630	510	
	7:00	2928	8840	LO	15.98	15.10	4.3	4.4	2271	2040	2015	2075	2143	1991	1885	1877	1935	1858	1811	1975	1945	2110	2330	2435	2462	490	
	8:00	2920	8861	LO	11.79	14.31	4.4	4.4	1848	1697	1668	1637	1588	1057	1436	1009	1249	1439	1467	1615	1606	1696	1723	1801	2279	350	
	9:00	2977	8874	HI	12.04	14.74	4.3	4.4	1881	1731	1661	1570	1509	1083	1408	1036	1420	1562	1457	1636	1625	1718	1740	1801	1852	140	
	10:00	2946	8887	HI	16.57	19.18	4.3	4.4	1895	1762	1698	1475	1447	1800	1476	1312	1402	1498	1412	1648	1636	1719	1741	1818	1869	190	
	11:00	2920	8919	LO	11.57	14.42	4.6	4.4	2311	2092	2003	2121	2096	2003	1908	1881	1931	1821	1788	1964	1942	2101	2238	2431	2497	450	
	12:00	2926	8945	HI	16.88	19.67	4.3	4.4	2548	2191	2044	2055	2124	2031	1927	1905	1993	1864	1821	2031	1974	2144	1875	1904	1965	380	
	1:00	2923	8971	HI	16.74	18.86	4.3	4.4	2386	2131	2062	2086	2152	2042	1957	1941	2007	1871	1811	2001	1950	2095	2309	2501	2622	470	
	2:00	2938	8992	HI	16.31	19.05	4.3	4.4	2441	2069	1981	1961	2072	1977	1890	1878	1976	1821	1724	1929	2055	2352	2522	2699	440		

\* 1.000 4 1.0000

TEST NO. 4

SYSTEM STATUS

PAGE 1 OF 1

DATE	TIME	ELAPSED TIME (HRS)	HYDROGEN OUTPUT METER	POWER INPUT		PRESSURE (PSIG)			COOLANT TEMP (°F)	FLOW RATE				GAS ANALYSIS			O <sub>2</sub> METER
				HIGH VOLTS	LOW VOLTS	MOD	PT	PO <sub>2</sub>		KOH	CMNT	N <sub>2</sub> H <sub>4</sub>	META	N <sub>2</sub>	CMNT	N <sub>2</sub>	
12-7	20:00	0	1463	28.7	20.8	36	4.7	116	1.9	5.9	3.8	.8	0	1.4	0	100	
12-7	20:00	2	1659	28.7	21.0	36	4.4	131	1.9	5.9	3.0	.8	0	2.0	0	120	
12-7	20:15					PO <sub>2</sub> = 140					137						
						Set to 150					Flow = 9.0						
						PO <sub>2</sub> control col					Set to 3.31						
						Note Prim to N <sub>2</sub> H <sub>4</sub> stat					Set to 2.24						
						Return to control to = PO <sub>2</sub>					Set to 1.57						
8:20						37	4.3	153	1.9	5.9	4.1						
8:35						Changed											
9:00						10 cc KOH (56)											
9:05						Removed											
9:10						Hi Sample Cathode											
9:30	4		1866	28.5	21.0	21.5	9.0	36	4.3	160	1.9	5.9	4.1	3.5	.8	8.5	
11:15						N <sub>2</sub> Sample Cathode											
11:30	6		2064	28.7	21.8	21.6	9.0	36	4.3	159	1.9	5.9	4.3	4.0	.8	8.5	
12-8	10:30	8	2121	28.5	21.5	28.5	9.5	31	4.3	156	1.9	5.9	4.25	3.3	.8	8.5	
3:18						0.1. Canceled											
3:32	10		2412	28.5	21.5	28.5	9.5	34	4.3	148	1.9	5.9	4.2	3.8	.8	8.5	
4:28						Removed N <sub>2</sub> H <sub>4</sub> & weight											
5:35	12																
5:32	12		2604	28.5	21.5	28.5	9.5	32	4.3	146	1.9	5.9	4.1	3.3	.8	8.5	
7:35	14		2935	28.3	21.4	28.3	9.4	29	4.4	154	1.9	5.9	4.5	3.3	.78	8.5	
8:30	15		-13	28.5	21.5	28.5	9.5	33	4.3	156	1.9	5.9	4.4	3.2	.78	8.5	
9:10	React					N <sub>2</sub> H <sub>4</sub> flow control valve to 5.93 turns											
9:30	16		131	28.5	21.8	28.5	9.5	35	4.3	156	1.9	5.9	4.4	3.2	.75	8.5	
9:45	Adjusted					N <sub>2</sub> makeup to 12.5 SS											
10:30	17		239	28.5	21.8	28.5	9.5	39	4.3	153	1.9	5.9	4.3	3.0	.75	8.5	
12:00	Removed					N <sub>2</sub> H <sub>4</sub> & weight											
1:30	20		524	28.5	21.4	28.5	9.5	39	4.3	143	1.9	5.9	4.2	3.0	.75	8.5	
3:00	coolant cycled off					for feed time in a long time											
4:30	23		771	28.4	21.5	28.4	9.5	37	4.3	153	1.9	5.9	4.5	3.0	.75	8.5	
5:00	24		Removed			10 cc KOH to 100											

TEST NO. - 4

SYSTEM STATUS

PAGE 3 OF —

DATE	TIME	ELAPSED TIME (HRS)	HYDROGEN OUTPUT METER	POWER INPUT		PRESSURE (PSIG)			COOLANT TEMP (°F)	FLOW RATE				GAS ANALYSIS									
				HIGH VOLTS	LOW VOLTS	MOD	PT	PO2		CAL	NH4	KCH	CHAMANT	N2 H2A	META	N2	BUB. SEC.	N2 - O2	CABIN	H2			
12-8	6:35	25 +	960	28.5	21.5	20.7	92	41	43	154	1.9	5.7	44	3.8	8.5	3.6	12.5	100	2700	2760 cc/min			
	9:40	27 +	1184		21.6	23	31	43	151	1.9	5.5	45	46	.8	2.0	3.6	12.4	100	2709	2080 cc/min			
	10:00	Adjusted high current on A & B down to ~ 13.5 amps																					
	10:35	Changed meta & N2 makeup to 20.55 and 4.835 respectively																					
	10:45	1383																100	27175				
	11:00	Note: Last cycle 440.7 cc N2H4 consumed																					
	11:10	297	N2 to 5.7 (440) Qmeta = 2.1 (945) 4 man size																				
	11:30	30	1440		28.5	9.1	35	42	147	1.9	5.9	50	52	-	4.8	2.1	5.8	-					
	11:45	flow check Qmeta = 9.54 cc/min (bevel)																					
12-9	1:34	32 +	1809		28.5	9.5	41	43	157	1.9	5.9	44.5	5.3	-	9.0	2.0	5.7		944 cc/min				
	2:00				Remade 10cc electrolyte																		
	3:30	34	1624		28.5	9.5	43	43	163	1.9	5.9	43.5	5.5	-	9.0	2.0	5.7		943 cc/min				
	5:33	36 +	1709		28.5	9.3	43.5	43	162	1.9	5.9	44.5	5.6	-	9.0	2.0	5.7		940 cc/min				
	7:45		2067		28.5	9.5	46	43	160	1.9	5.9	50	5.6		9.0	2.1	5.7		935 cc/min	adj to 946			
	9:10	adjusted meta to 990 cc/min (Current not cycling)																					
	9:30	40																	999 cc/min				
	9:40		2236		28.5	9.5	42	44	160	1.9	5.9	49	5.8	.8	8.4	2.1	5.7	100	955 cc/min	(12749 am)			
	10:30	41	2307		28.5	9.5	38	43	160	1.9	5.9	48	6.3	-	8.9	2.1	5.7	100	955 cc/min	2741 10 cc			
	1:30	44	2541		28.5	9.5	37	44	157	1.9	5.9	50	5.8	-	8.6	2.1	5.7	100	950 cc/min	2751 16.15			
	2:30	45	2632		28.5	9.5	45	43	156	1.9	5.9	51	6.5	.8	8.6	2.1	5.7	100		2759 20 cc 10:30			
	2:45	Changed meta & N2 makeup to 5-man rate																					
	3:45																						
	4:00	Adjusted N2 makeup to 6.655																					
	4:35	47 +	2447		28.5	9.4	45	44	155	1.9	5.9	49	6.0	-	8.5	2.3	6.6	100	1055 cc/min	2754 10 cc 11:30 30 cc 1:30			
	4:55	Changed meta & N2 makeup to 6-man rate																					
	6:50	Drain N2H4 & re-charge 12.98.9 used																					
	7:10	7:35 Adj Start set to 10 man level																	(11085) Qmeta = 31	1830 70 18 30			
	7:40	50 +	0111		28.5	9.1	31	43	156	1.9	6.0	47	6.7	-	9.0	3.3	8.1	-	2763	cc/min			
	9:30	52	0309		28.7	9.1	37	43	155	1.9	5.8	47	7.2	.8	8.5	3.3	11.0	-	2771				
	11:00	54	0549		28.0	21.8	34	43	152	1.9	5.5	46	6.0	.8	9.0	3.3	11.0	-	2779				
12-10	11:16				ANALYST	14E-78	FACE	22	36.55	=	214.5 cc/min	(12 man rate)											

WET TEST 82



TEST ND. 4

SYSTEM STATUS

DATE	TIME	ELAPSED TIME (HRS)	HYDROGEN OUTPUT METER	POWER INPUT		PRESSURE (PSIG)				COOLANT TEMP (°F)	FLOW RATE			GAS ANALYSIS			
				HIGH VOLTS	LOW VOLTS	MOD	PT	POS	CAL		NH4	KOH	COUMT/N4	WETA	N2	N2-O2	CABIN
12-21-92	0719	57		28.5	22.5	32	4.5	146	1.9	5.9	43.0	70.8	—	36	110		
	1:52			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	0910	59		28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	1000	61		28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	1321	63		28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	8:08			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	9:15			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	9:30			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	10:30			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	11:30			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	1:30			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	3:45			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	4:30			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	6:30			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	7:30			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	8:30			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	8:30			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	8:30			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	8:35			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	9:00			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	9:10			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	9:30			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	10:00			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	11:30			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	1:30			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	3:32			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	4:31			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	5:45			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	7:34			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		
	8:00			28.5	22.5	32	4.5	146	1.9	5.9	43.5	70.8	—	36	110		

H<sub>2</sub>O leak  
.57 min  
10 cm  
trans



TEST NO. 4

## MODULE STATUS

PAGE 2 OF

DATE	TIME	HIGH TIMER	LOW TIMER	MODE	MOD VOLTS		MOD AMPS		CELL VOLTAGE												EFFLUENT POL	NZHA CONC., *					
					A	B	A	B	1A	2A	3A	4A	5A	6A	7A	8A	9A	10A	11A	12A			13A	14A			
12-5	4:30	45:37	10:49	LO	14.07	16.14	6.7	6.8	1805	1851	1764	1722	1777	1712	1701	1670	1674	1722	1676	1810	1780	1815	1829	1892	1925	290	1050 3:45 AM
30	5:30	45:37	11:49	LO	14.03	16.05	6.7	6.8	1805	1851	1764	1719	1771	1710	1703	1668	1672	1718	1673	1801	1766	1788	1805	1862	1898	270	1050 4:55 AM
37	6:30	45:37	12:49	LO	14.00	16.00	6.7	6.8	1805	1851	1764	1717	1777	1717	1714	1673	1677	1722	1677	1803	1770	1793	1812	1873	1909	265	5:45 1000
38	7:40	45:37	13:46	LO	14.02	16.01	6.7	6.8	1805	1851	1764	1711	1775	1716	1707	1671	1676	1721	1675	1802	1767	1797	1819	1885	1933	250	6:45 1000
40	8:45	45:37	14:55	LO	14.04	16.05	6.7	6.8	1805	1851	1764	1710	1773	1713	1703	1669	1670	1721	1673	1802	1771	1802	1825	1894	1948	240	8:00-1000
41	10:30	45:59	16:27	H1	16.06	18.91	13.4	13.4	2197	2230	1947	1940	2045	1919	1888	1850	1857	1927	1887	2118	2095	2095	2095	2095	2095	380	11:30
44	11:30	45:37	16:27	LO	14.06	16.18	6.7	6.8	1805	1851	1764	1711	1773	1713	1703	1669	1670	1721	1673	1802	1771	1802	1825	1894	1932	240	
45	2:30	45:37	16:91	LO	14.13	16.17	6.7	6.8	1805	1851	1764	1711	1776	1712	1704	1671	1677	1730	1678	1813	1774	1801	1831	1876	1926	220	
47	4:30	45:39	17:09	LO	14.08	16.19	6.7	6.8	1805	1851	1764	1711	1776	1712	1704	1671	1677	1730	1678	1813	1774	1801	1831	1876	1926	220	
48	6:35	46:44	17:41	H1	16.62	17.75	13.8	13.8	2335	2433	2029	1954	2155	1951	1918	1861	1866	1927	1887	2119	2163	2163	2163	2163	2163	350/220	
50	7:35	46:44	17:41	H1	16.67	17.75	13.8	13.8	2335	2433	2029	1954	2155	1951	1918	1861	1866	1927	1887	2119	2163	2163	2163	2163	2163	270/410	
51	8:30	47:05	18:19	H1	16.91	17.77	14.0	14.0	2335	2433	2029	1954	2155	1951	1918	1861	1866	1927	1887	2119	2163	2163	2163	2163	2163	250/390	
52	9:30	47:35	19:01	LO	14.11	16.31	6.7	6.8	1805	1851	1764	1714	1776	1709	1694	1660	1659	1734	1701	1825	1802	1834	1867	1912	200/370		
53	10:30	48:05	19:17	H1	13.92	16.46	6.9	6.9	1337	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362		
54	11:30	48:05	20:17	LO	14.11	16.21	6.9	6.9	1337	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	180/360	
55	12:30	49:24	20:44	LO	14.12	16.19	6.9	6.9	1337	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	1362	160/350	
56	1:30	49:34	20:54	LO	16.01	19.13	13.7	13.7	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	1940	
57	2:30	50:01	20:58	H1	16.54	18.96	13.6	13.6	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	320/41	1050 1:25 PM
58	3:30	50:37	21:05	H1	17.41	20.24	14.0	14.0	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	310	
59	4:30	51:46	21:52	H1	17.17	19.94	14.0	14.0	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	140	1050 4:20 PM
60	5:30	51:57	21:59	H1	17.13	19.98	14.0	14.0	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	450	
61	6:30	52:31	22:10	H1	16.95	18.93	13.9	13.9	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	420	
62	7:30	52:46	22:59	LO	14.04	16.44	6.7	6.8	1805	1851	1764	1724	1787	1701	1678	1666	1677	1737	1694	1823	1798	1828	1851	1902	1938	245	
63	8:30	53:57	23:50	H1	16.66	19.46	13.2	13.2	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	210	
64	9:30	54:57	24:42	LO	14.00	16.30	6.7	6.8	1805	1851	1764	1724	1787	1701	1678	1666	1677	1737	1694	1823	1798	1828	1851	1902	1938	380	
65	10:30	55:55	25:42	H1	16.36	19.05	13.2	13.2	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	2351	440	
66	11:30	56:42	26:47	LO	14.41	16.52	7.3	7.3	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	430	
67	12:30	57:45	27:36	LO	14.62	16.60	7.4	7.4	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	250	
68	1:30	58:45	28:31	LO	14.67	16.74	7.4	7.4	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	310/410	1050 4:20
69	2:30	59:45	29:28	LO	14.62	16.74	7.4	7.4	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	400/460	
70	3:30	60:45	30:28	LO	14.62	16.74	7.4	7.4	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	1952	460/490	

\* EVERY 4 HOURS

TEST NO. 4

MODULE STATICS

PAGE 3 OF

DATE	TIME	HIGH TIMER	LOW TIMER	MODE	MOD VOLTS		MOD AMPS		CELL VOLTAGE												EFFLUENT POL	N2H4 CONC. *					
					A	B	A	B	1A	2A	3A	4A	5A	6A	7A	8A	9B	8B	7B	6B			5B	4B	3B	2B	1B
12-10	8:30 AM	57.18	21.58	LI	16.62	17.24	12.01	12.01	2.27	2.27	2.09	2.05	2.26	2.26	1.93	1.93	1.90	1.92	1.90	2.20	2.05	2.16	2.34	2.42	2.54	540/520	
	9:30	58.07	21.59	LO	14.71	16.53	12.01	12.01	1.90	1.86	1.83	1.85	2.05	2.05	1.79	1.78	1.70	1.73	1.70	1.84	1.81	1.90	1.90	1.90	1.92	500/520	
	10:30	58.64	28.25	LO	14.80	16.62	12.01	12.01	1.94	1.88	1.87	1.85	2.05	2.05	1.73	1.70	1.75	1.73	1.74	1.84	1.81	1.93	1.93	1.94	1.96	480/420	
	11:30	57.30	21.56	LI	16.71	17.58	12.01	12.01	2.27	2.27	2.09	2.03	2.26	2.26	1.93	1.93	1.90	1.92	1.90	2.20	2.05	2.16	2.34	2.42	2.54	370/440	
	12:30	59.70	28.73	LO	14.38	16.31	12.01	12.01	1.90	1.86	1.83	1.85	2.05	2.05	1.79	1.78	1.70	1.73	1.70	1.84	1.81	1.90	1.90	1.90	1.92	410	
	1:30	60.76	28.80	LI	16.53	19.91	12.01	12.01	2.27	2.27	2.09	2.03	2.26	2.26	1.93	1.93	1.90	1.92	1.90	2.20	2.05	2.16	2.34	2.42	2.54	365	
	2:30	61.49	29.11	LO	14.22	16.61	12.01	12.01	1.90	1.86	1.83	1.85	2.05	2.05	1.79	1.78	1.70	1.73	1.70	1.84	1.81	1.90	1.90	1.90	1.92	230	
	3:30			LO	14.19	16.96	12.01	12.01	1.90	1.86	1.83	1.85	2.05	2.05	1.79	1.78	1.70	1.73	1.70	1.84	1.81	1.90	1.90	1.90	1.92	210	
	4:30	62.81	29.76	LO	15.60	18.15	12.01	12.01	2.17	2.05	1.94	1.90	2.15	2.15	1.98	1.94	1.82	1.89	1.80	2.05	2.05	2.05	2.13	2.19	2.28	345	
	5:30	63.27	30.05	LI	15.11	18.05	12.01	12.01	2.18	1.98	1.94	1.90	1.95	1.95	1.80	1.77	1.74	1.85	1.82	2.08	2.05	2.13	2.27	2.28	2.33	350	304M
	6:30	63.75	30.59	LO	14.12	16.54	12.01	12.01	1.87	1.80	1.76	1.74	1.85	1.85	1.71	1.68	1.70	1.74	1.72	1.84	1.83	1.86	1.88	1.88	1.88	380	
	7:30	64.18	30.67	LO	14.26	16.94	12.01	12.01	1.90	1.87	1.80	1.76	1.87	1.87	1.71	1.69	1.70	1.74	1.71	1.81	1.78	1.80	1.80	1.80	1.80	470	

1012  
5:10 AM